

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 17:45:55 ; Search time 1500.84 Seconds
(without alignments)
532.600 Million cell updates/sec

Title: US-10-729-421-40

Perfect score: 21

Sequence: 1 cagtgcagtcaggtctagct 21

Scoring table: IDENTITY NUC

Gapop 10'0 , Gapext 1.0

Searched: 34239544 segs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database :

EST:*

1: gb_est1:*

2: gb_est2:*

3: gb_hc:*

4: gb_est3:*

5: gb_est4:*

6: gb_est5:*

7: gb_est6:*

8: gb_ges1:*

9: gb_ges2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----------|--------------------|
| 1 | 19 | 90.5 | 1041 | CNS04CAJ | AL284212 Tetraodon |
| 2 | 18.4 | 87.6 | 1157 | CD500170 | CD500170 CDA43-E06 |
| 3 | 17.8 | 84.8 | 211 | A2995872 | A2995872 2M0281B23 |
| C 4 | 17.8 | 84.8 | 611 | A2218630 | A2218630 Sheared D |
| 5 | 17.8 | 84.8 | 878 | CR794665 | CR794665 GROAA12A |
| C 6 | 17.4 | 84.8 | 4261 | AK083880 | AK083880 Mus muscu |
| C 7 | 17.4 | 82.9 | 380 | CL289600 | CL289600 ZMMBB064 |
| 8 | 17.4 | 82.9 | 718 | CG176005 | CG176005 PUTMY95TD |
| C 9 | 17.4 | 82.9 | 722 | BF607360 | BF607360 MYL 00030 |
| C 10 | 17.4 | 82.9 | 772 | CC340825 | CC340825 OGQAQ84TH |
| C 11 | 17.4 | 82.9 | 772 | CG211386 | CG211386 OGSCS03TC |
| C 12 | 17.4 | 82.9 | 805 | CG362883 | CG362883 OGYSB24TH |
| C 13 | 17.4 | 82.9 | 816 | B1548151 | B1548151 603189492 |
| C 14 | 17.4 | 82.9 | 817 | CC340835 | CC340835 OGQAQ84TV |
| C 15 | 17.4 | 82.9 | 923 | CG211861 | CG211861 OGLAW29TV |
| 16 | 17 | 81.0 | 382 | BY609340 | BY609340 BY609340 |
| 17 | 17 | 81.0 | 408 | BY649695 | BY649695 BY649695 |
| C 18 | 17 | 81.0 | 622 | CA124343 | CA124343 SCQGLR108 |
| C 19 | 17 | 81.0 | 633 | AZ069535 | AZ069535 RPCI-23-4 |
| C 20 | 17 | 81.0 | 642 | B8650662 | B8650662 BB650662 |
| C 21 | 17 | 81.0 | 655 | BB293162 | BB293162 BB293162 |
| 22 | 17 | 81.0 | 768 | AG457137 | AG457137 Mus muscu |
| 23 | 17 | 81.0 | 781 | AG557182 | AG557182 Mus muscu |
| 24 | 17 | 81.0 | 949 | CF411086 | CF411086 CH3#071_D |

| | | | | | | |
|------|------|------|------|---|----------|--------------------|
| C 25 | 17 | 81.0 | 2445 | 3 | AK044974 | AK044974 Mus muscu |
| C 26 | 17 | 81.0 | 2466 | 3 | AK013040 | AK013040 Mus muscu |
| C 27 | 17 | 81.0 | 2559 | 3 | AK017012 | AK017012 Mus muscu |
| C 28 | 17 | 81.0 | 3256 | 3 | AK082079 | AK082079 Mus muscu |
| C 29 | 17 | 81.0 | 3417 | 3 | AK081942 | AK081942 Mus muscu |
| C 30 | 16.8 | 80.0 | 306 | 2 | AW846933 | AW846933 RC3-CT019 |
| 31 | 16.8 | 80.0 | 330 | 1 | AU249500 | AU249500 AU249500 |
| 32 | 16.8 | 80.0 | 335 | 1 | AA050319 | AA050319 mjl4a05.r |
| 33 | 16.8 | 80.0 | 387 | 5 | BY077663 | BY077663 BY077663 |
| 34 | 16.8 | 80.0 | 400 | 1 | AA004028 | AA004028 mg80g02.r |
| 35 | 16.8 | 80.0 | 400 | 1 | AA117056 | AA117056 mn29c06.r |
| 36 | 16.8 | 80.0 | 407 | 1 | AA003782 | AA003782 mg62e10.r |
| 37 | 16.8 | 80.0 | 407 | 2 | BE656207 | BE656207 UI-M-EH0- |
| 38 | 16.8 | 80.0 | 414 | 1 | AA052553 | AA052553 mc66d04.r |
| 39 | 16.8 | 80.0 | 419 | 7 | W79973 | W79973 me90609.r1 |
| 40 | 16.8 | 80.0 | 426 | 1 | AA008849 | AA008849 mg98f08.r |
| 41 | 16.8 | 80.0 | 435 | 5 | BY273688 | BY273688 BY273688 |
| 42 | 16.8 | 80.0 | 436 | 5 | BY240303 | BY240303 BY240303 |
| 43 | 16.8 | 80.0 | 442 | 5 | BY051479 | BY051479 BY051479 |
| 44 | 16.8 | 80.0 | 443 | 6 | CB788954 | CB788954 AMGNNUC:M |
| 45 | 16.8 | 80.0 | 461 | 5 | EX529253 | EX529253 BX529253 |
| 46 | 16.8 | 80.0 | 466 | 7 | W89580 | W89580 mf73f08.r1 |
| 47 | 16.8 | 80.0 | 467 | 1 | AJ647590 | AJ647590 AJ647590 |
| 48 | 16.8 | 80.0 | 467 | 2 | BB863212 | BB863212 BB863212 |
| 49 | 16.8 | 80.0 | 469 | 1 | AJ647892 | AJ647892 AJ647892 |
| 50 | 16.8 | 80.0 | 508 | 6 | CA533118 | CA533118 C0345E08- |
| 51 | 16.8 | 80.0 | 510 | 7 | W89380 | W89380 mf73g08.r1 |
| 52 | 16.8 | 80.0 | 512 | 2 | BF556581 | BF556581 UI-R-E1-f |
| 53 | 16.8 | 80.0 | 515 | 2 | BE374887 | BE374887 601226811 |
| 54 | 16.8 | 80.0 | 523 | 8 | BZ132704 | BZ132704 CH230-481 |
| 55 | 16.8 | 80.0 | 542 | 1 | AA712019 | AA712019 ui60d05.r |
| 56 | 16.8 | 80.0 | 544 | 7 | CF732719 | CF732719 UI-M-HA0- |
| 57 | 16.8 | 80.0 | 546 | 8 | BH347218 | BH347218 CH230-42D |
| 58 | 16.8 | 80.0 | 549 | 4 | BG276278 | BG276278 uv02e10.y |
| 59 | 16.8 | 80.0 | 552 | 1 | AA682125 | AA682125 vj13b03.r |
| 60 | 16.8 | 80.0 | 561 | 2 | AA530593 | AA530593 vj49g09.r |
| 61 | 16.8 | 80.0 | 568 | 1 | BF452552 | BF452552 mago1d05. |
| 62 | 16.8 | 80.0 | 570 | 1 | AV597290 | AV597290 AV597290 |
| 63 | 16.8 | 80.0 | 576 | 1 | AV595391 | AV595391 AV595391 |
| 64 | 16.8 | 80.0 | 582 | 1 | AL792667 | AL792667 AL792667 |
| 65 | 16.8 | 80.0 | 584 | 5 | BQ840540 | BQ840540 mah69f08. |
| 66 | 16.8 | 80.0 | 588 | 9 | CG672097 | CG672097 RRM266 Ba |
| 67 | 16.8 | 80.0 | 596 | 4 | BG100744 | BG100744 uy14c01.y |
| 68 | 16.8 | 80.0 | 600 | 4 | B1985696 | B1985696 3144-63 M |
| 69 | 16.8 | 80.0 | 617 | 2 | AW412020 | AW412020 uo5sh02.y |
| 70 | 16.8 | 80.0 | 619 | 7 | CR421241 | CR421241 CR421241 |
| 71 | 16.8 | 80.0 | 624 | 7 | CK621976 | CK621976 ml31a12.y |
| 72 | 16.8 | 80.0 | 626 | 4 | BJ774417 | BJ774417 BJ774417 |
| 73 | 16.8 | 80.0 | 627 | 7 | CN119882 | CN119882 ECOCNA002 |
| 74 | 16.8 | 80.0 | 637 | 4 | BG099869 | BG099869 uy13c02.y |
| 75 | 16.8 | 80.0 | 641 | 6 | CD766336 | CD766336 AGENCOURT |
| 76 | 16.8 | 80.0 | 642 | 1 | AL863627 | AL863627 AL863627 |
| 77 | 16.8 | 80.0 | 642 | 1 | AL878478 | AL878478 AL878478 |
| 78 | 16.8 | 80.0 | 649 | 7 | CF732690 | CF732690 UI-M-HA0- |
| 79 | 16.8 | 80.0 | 651 | 5 | EX315194 | EX315194 BX315194 |
| 80 | 16.8 | 80.0 | 655 | 6 | CD772714 | CD772714 AGENCOURT |
| C 81 | 16.8 | 80.0 | 660 | 8 | AZ428022 | AZ428022 IM0210N17 |
| C 82 | 16.8 | 80.0 | 668 | 4 | BG381620 | BG381620 UI-R-CT0- |
| C 83 | 16.8 | 80.0 | 685 | 9 | AG151855 | AG151855 Pan trogl |
| 84 | 16.8 | 80.0 | 688 | 5 | BQ746626 | BQ746626 UI-M-ER0- |
| 85 | 16.8 | 80.0 | 701 | 7 | CK654484 | CK654484 AGENCOURT |
| 86 | 16.8 | 80.0 | 720 | 8 | AZ247538 | AZ247538 RPCI-23-9 |
| C 87 | 16.8 | 80.0 | 720 | 8 | AZ247542 | AZ247542 RPCI-23-9 |
| C 88 | 16.8 | 80.0 | 734 | 7 | CF148724 | CF148724 AGENCOURT |
| 89 | 16.8 | 80.0 | 734 | 7 | CO806312 | CO806312 AGENCOURT |
| 90 | 16.8 | 80.0 | 738 | 6 | CB598900 | CB598900 AGENCOURT |
| C 91 | 16.8 | 80.0 | 744 | 9 | AG527299 | AG527299 Mus muscu |
| C 92 | 16.8 | 80.0 | 749 | 7 | CO562201 | CO562201 AGENCOURT |
| 93 | 16.8 | 80.0 | 753 | 7 | CF411613 | CF411613 CR441613 |
| 94 | 16.8 | 80.0 | 765 | 7 | CF149901 | CF149901 AGENCOURT |
| 95 | 16.8 | 80.0 | 768 | 9 | CC553492 | CC553492 CH240_459 |
| 96 | 16.8 | 80.0 | 822 | 9 | AY413996 | AY413996 Mus muscu |
| 97 | 16.8 | 80.0 | 829 | 5 | B0412600 | B0412600 602954786 |

98 16.8 80.0 835 4 BI684946
 c 99 16.8 80.0 844 9 CR017849 Forward s
 100 16.8 80.0 847 7 CK793898 AGENCOURT

ALIGNMENTS

RESULT 1
 CDS04CAJ
 LOCUS
 DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone
 099K23 of library G from Tetraodon nigroviridis, genomic survey
 sequence.

ACCESSION AL284212
 VERSION AL284212.1 GI:8022590
 KEYWORDS GSS; genome survey sequence.
 SOURCE Tetraodon nigroviridis
 ORGANISM Tetraodon nigroviridis
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
 Tetraodontoidea; Tetraodontidae; Tetraodon.

REFERENCE 1
 AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
 Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
 Saurin,W. and Weissenbach,J.

TITLE Estimate of human gene number provided by genome-wide analysis
 using Tetraodon nigroviridis DNA sequence
 JOURNAL Nat. Genet. 25 (2), 235-238 (2000)
 MEDLINE 20296633
 PUBMED 10835645

REFERENCE 2
 AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Ozouf-Costaz,C.,
 Fizames,C., Fischer,C., Bouneau,L., Billault,A., Quetier,F.,
 Saurin,W., Bernot,A. and Weissenbach,J.

TITLE Characterization and repeat analysis of the compact genome of the
 freshwater pufferfish Tetraodon nigroviridis

JOURNAL Genome Res. 10 (7), 939-949 (2000)
 MEDLINE 20359837
 PUBMED 10899143

REFERENCE 3 (bases 1 to 1041)
 AUTHORS Genoscope.
 TITLE Direct Submission

JOURNAL Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :
 BP 191 91006 EVRY cedex - FRANCE (E-mail : seqrefgenoscope.cns.fr
 - Web : www.genoscope.cns.fr)

COMMENT This sequence is a single read and was generated as part of a large
 scale clone-end sequencing project of the Tetraodon nigroviridis
 genome. For more information, please take a look at
 http://www.genoscope.cns.fr/Tetraodon.

FEATURES

source

Location/Qualifiers
 1..1041
 /organism="Tetraodon nigroviridis"
 /mol_type="genomic DNA"
 /db_xref="taxon:99883"
 /clone="099K23"
 /clone_lib="G"
 /note="Genoscope sequence ID : C08G099AF12LP1-end : T7"

ORIGIN

Query Match 90.5%; Score 19; DB 9; Length 1041;
 Best Local Similarity 90.5%; Pred. No. 2e+02;
 Matches 19; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTCTAGCT 21
 :|||||
 Db 942 SAGTGACATGCAGGCTCTACCT 962

RESULT 2
 CDS00170
 LOCUS

DEFINITION CDA43-E06.xid-t SHGC-CDA Gasterosteus aculeatus cDNA clone
 CDA43-E06 5', mRNA sequence.
 ACCESSION CD500170
 VERSION CD500170.1 GI:31427201
 KEYWORDS EST.
 SOURCE Gasterosteus aculeatus (three spined stickleback)
 ORGANISM Gasterosteus aculeatus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 Acanthomorpha; Acanthopterygii; Percomorpha; Gasterosteiformes;
 Gasterosteidae; Gasterosteus.

REFERENCE 1 (bases 1 to 1157)
 AUTHORS Kingsley,D.M., Peichel,C., Balabhadra,S., Grinwood,J., Dickson,M.,
 Schmutz,J. and Myers,R.M.

TITLE Expressed sequence tags from Gasterosteus aculeatus
 JOURNAL Unpublished (2003)
 COMMENT Contact: Kingsley, DM
 HHMI and Department of Developmental Biology
 Stanford University School of Medicine
 Beckman Center B300, 279 Campus Drive, Stanford, CA 94305-5329, USA
 Tel: 650 725 5954
 Fax: 650 725 7739
 Email: kingsley@cgm.stanford.edu
 Plate: 43

High quality sequence stop: 785.

FEATURES

source

Location/Qualifiers
 1..1157
 /organism="Gasterosteus aculeatus"
 /mol_type="mRNA"
 /strain="Salinas river, CA"
 /db_xref="taxon:69293"
 /clone="CDA43-E06"
 /sex="mixed male and female"
 /tissue_type="heads and internal organs combined"
 /dev_stage="adult"
 /clone_lib="SHGC-CDA"
 /note="Vector: lambda ZAP Express/pBK-CMV; Site 1: EcoRI
 (5' adaptor); Site 2: XhoI (3' linker primer); The mixed
 organ cDNA library was generated using the ZAP-cDNA method
 by Stratagene. First strand cDNA synthesis was primed with
 a 50 bp linker primer containing an oligo dt sequence
 preceded by a synthetic XhoI site. 5 prime adaptors were
 used containing an EcoRI cohesive end. The finished cDNAs
 were inserted in to the ZAP express vector
 unidirectionally in the sense orientation with respect to
 the lacZ promoter of pBK-CMV. An amplified library was
 prepared from approximately 3 million primary clones in
 the lambda ZAP Express vector. In vivo excision was then
 used to generate individual pBK-CMV phagemid clones for
 EST sequencing."

ORIGIN

Query Match 87.6%; Score 18.4; DB 6; Length 1157;
 Best Local Similarity 95.0%; Pred. No. 4e+02;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 2 AGTGACATGCAGGCTCTAGCT 21
 :|||||
 Db 132 AGTGACATGCAGGCTCTACCT 151

RESULT 3

AZ995872

LOCUS

DEFINITION AZ995872 211 bp DNA linear GSS 27-APR-2001
 clone UUGC2M0281B23 R, genomic survey sequence.

ACCESSION

AZ995872

VERSION

AZ995872.1

KEYWORDS

GSS.

SOURCE

Mus musculus (house mouse)

ORGANISM

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 211)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Iqbal,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0281 row: B column: 23
 Seq primer: CACACGGAACACGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 211.

FEATURES
 source
 1..211
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC2M0281B23"
 /sex="Female"
 /lab_host="E. coli strain XL10-Gold, Tl-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC2M library"
 /notes="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (female) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pW42 (gi|4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN

Query Match 84.8%; Score 17.8; DB 8; Length 211;
 Best Local Similarity 90.5%; Pred. No. 6.1e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTCTAGCT 21
 |||||
 Db 136 CAGAGCCATGCAGGCTCTAGCT 156

RESULT 4
 LOCUS AZ218630/c 611 bp DNA linear GSS 09-JUN-2000
 DEFINITION Sheared DNA-82B8.TP Sheared DNA Trypanosoma brucei genomic clone
 Sheared DNA-82B8, genomic survey sequence.
 ACCESSION AZ218630
 VERSION AZ218630.1 GI:8436430
 KEYWORDS GSS.
 SOURCE Trypanosoma brucei
 ORGANISM Trypanosoma brucei
 Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma.

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 COMMENT

1 (bases 1 to 611)
 El-Sayed,N., Zhao,S., Zhao,H., Gill,S., Suh,E., Malek,J., Fujii,C., Gerrard,C., Leech,V., de Jong,P., Ullu,E., Melville,S., Donelson,J., Fraser,C. and Adams,M.
 Determination of clone end sequences from Trypanosoma brucei GUTat 10.1 sheared DNA library
 Unpublished (1999)
 Other GSSs: Sheared DNA-82B8.TR
 Contact: Najib M. El-Sayed
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: nelsayed@tigr.org
 Clones are derived from the Trypanosoma brucei GUTat 10.1 sheared DNA library constructed at TIGR. Clones will be available for distribution through Research Genetics, Alabama, USA. Sheared DNA end sequences search page: <http://www.tigr.org/cdb/mdb/tbdb/>.
 Seq primer: M13-Forward
 Class: shotgun.

FEATURES
 Location/Qualifiers
 1..611
 /organism="Trypanosoma brucei"
 /mol_type="genomic DNA"
 /strain="TREU927/4 GUTat 10.1"
 /db_xref="taxon:5691"
 /clone="Sheared DNA-82B8"
 /clone_lib="Sheared DNA"
 /notes="Vector: pUC18; Site 1: SmaI; Constructed at The Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (approx 2 kb). The v + i method used for the library construction is described in detail in Smith, H.O. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaudin and B. Borell, Oxford University Press, 1999)."

ORIGIN

Query Match 84.8%; Score 17.8; DB 8; Length 611;
 Best Local Similarity 90.5%; Pred. No. 7.2e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTCTAGCT 21
 |||||
 Db 548 CAGTGGCATGCAGGCTCTAGTT 528

RESULT 5

LOCUS CR794665 878 bp DNA linear GSS 24-SEP-2004
 DEFINITION GR05AA12AC07RM1 INRA BAC Bos taurus genomic clone INRAB_225E12, DNA sequence, genomic survey sequence.

ACCESSION CR794665
 VERSION CR794665.1 GI:52675664
 KEYWORDS GSS.
 SOURCE Bos taurus (cow)
 ORGANISM Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovinae; Bos.

REFERENCE 1 (bases 1 to 878)
 AUTHORS Eggen,A., Schibler,L. and Roy,A.
 TITLE Bovine BAC End Sequences from the INRA bovine BAC library
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 878)
 Genoscope.

Direct Submission
 TITLE Submitted (20-SEP-2004) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)

```

- Web : www.genoscope.cns.fr)
Contact: Andre Eggen
Department of Animal Genetics - LGbc
INRA
78350 Jouy-en-Josas, France
Tel: 33 1 34 65 24 24
Fax: 33 1 34 65 24 78
Email: eggen@jouy.inra.fr
Clones are derived from the INRA bovine BAC library
(http://locus.jouy.inra.fr/fpc/cattle_bac_map.htm). For BAC library
availability, please contact Andre Eggen (eggen@jouy.inra.fr). This
work was undertaken as part of the International Bovine BAC
Mapping Consortium (IBBMC) by INRA (Jouy-en-Josas) and Genoscope
(Evry) primer: 225 row: E column: 12
Seq primer: M13 Reverse
Class: BAC ends.
FEATURES
    source
        location/Qualifiers
            1..878
            /organism="Bos taurus"
            /mol_type="genomic DNA"
            /strain="breed: Holstein"
            /db_xref="taxon:9913"
            /clone="INRAB_225E12"
            /sex="Male"
            /cell_type="fibroblast"
            /clone_lib="INRA bovine BAC"
            /note="Vector: pBeloBAC11; Site 1: HindIII; Holstein bull;
            INRA Bovine BAC library (Male) produced by Andre
            Eggen-Genoscope sequence ID : GR0AAA12AC07RM1"
ORIGIN
    Query Match      84.8%; Score 17.8; DB 9; Length 878;
    Best Local Similarity 90.5%; Pred No. 7;e+02;
    Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCCTAGCT 21
    |||||
Db 297 CAGTCACAAGCAGGTCCTAGCT 317

RESULT 6
AK083880/c
LOCUS
DEFINITION Mus musculus 12 days embryo spinal ganglion cDNA, RIKEN full-length
enriched library, clone:DJ30043C18 product:unclassified, full
insert sequence.
ACCESSION AK083880
VERSION AK083880.1 GI:26101557
KEYWORDS HTC; CAP trapper.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1
REFERENCE
AUTHORS Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K.,
Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
TITLE High-efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 99279253
PUBMED 10349636
2
REFERENCE
AUTHORS Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K.,
Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
TITLE Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE 20499374
PUBMED 11042159
3
REFERENCE
AUTHORS Shibata,K., Itoh,M., Aizawa,K., Nagaoaka,S., Sasaki,N., Carninci,P.,
Konno,H., Akiyama,J., Nishi,K., Kitsuunai,T., Tashiro,H., Itoh,M.,
Sund,N., Ishii,Y., Nakamura,S., Hazama,M., Nishine,T., Harada,A.,
Yamamoto,R., Matsumoto,H., Sakaguchi,S., Ikegami,T., Kashiwagi,K.,
Fujiwaki,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E., Watahiki,M.,
Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsuura,S., Kawai,J.,
Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and Hayashizaki,Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer
JOURNAL Genome Res. 10 (11), 1757-1771 (2000)
PUBMED 11076861
4
REFERENCE
AUTHORS The RIKEN Genome Exploration Research Group Phase II Team and the
PANTOM Consortium.
TITLE Functional annotation of a full-length mouse cDNA collection
JOURNAL Nature 409, 685-690 (2001)
REFERENCE
AUTHORS The PANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
TITLE Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
JOURNAL Nature 420, 563-573 (2002)
REFERENCE
AUTHORS Adachi,J., Aizawa,K., Akimura,T., Arakawa,T., Bono,H., Carninci,P.,
Fukuda,S., Furuno,M., Hanagaki,T., Hara,A., Hashizume,W.,
Hayashida,K., Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,T.,
Hori,F., Imotani,K., Ishii,Y., Itoh,M., Kagawa,I., Kasukawa,T.,
Kato,H., Kawai,J., Kojima,Y., Kondo,S., Konno,H., Kouda,M.,
Koya,S., Kurihara,C., Matsuyama,T., Miyazaki,A., Murata,M.,
Nakamura,M., Nishi,K., Nomura,K., Numazaki,R., Ohno,M., Ohsato,N.,
Okazaki,Y., Saito,R., Saitoh,H., Sakai,C., Sakai,K., Sakazume,N.,
Sogabe,Y., Tagami,M., Tagawa,A., Takahashi,F., Takaku-Akahira,S.,
Takeda,Y., Tanaka,T., Tomaru,A., Toya,T., Yasunishi,A.,
Muramatsu,M. and Hayashizaki,Y.
Direct Submission
Submitted (16-APR-2002) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.jp,
URL:http://genome.gsc.riken.jp/, Tel:81-45-503-9222,
Fax:81-45-503-9216)
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Please visit our web site for further details.
URL:http://genome.gsc.riken.jp/
URL:http://fantom.gsc.riken.jp/.
FEATURES
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            /strain="C57BL/6J"
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            /db_xref="taxon:10090"
            /clone="DJ30043C18"
            /clone_lib="RIKEN full-length enriched mouse cDNA library"
            /dev_stage="12 days embryo"
            /note="unclassified"
            misc_feature
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                /note="unclassified"
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    Best Local Similarity 90.5%; Pred No. 1e+03;
    Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCCTAGCT 21
    |||||
Db 1497 CAGTGATATCGAGGTGTAGCT 1477

RESULT 7
CL289600/c

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LOCUS Cl289600 380 bp DNA linear GSS 10-FEB-2004
 DEFINITION ZMMBB0641H10r ZMMBBB (HindIII) Zea mays genomic clone
 VERSION ZMMBB0641H10 3', genomic survey sequence.
 KEYWORDS Cl289600 Cl289600.1 GI:42503987
 SOURCE GSS.
 ORGANISM Zea mays
 Zea mays
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.
 1 (bases 1 to 380)
 Bhatti,A.K., Young,S., Kaychok,S., Keizer,G., Bronzino,A.C.,
 Zohovetz,V., Fuks,Y., Yu,Y., Wing,R. and Messing,J.
 Sequencing of the maize genome at PGIR (2003c)
 TITLE Unpublished (2003)
 JOURNAL
 COMMENT Contact: Bhatti,A.K.
 Dr.Joschim Messing's lab
 The Plant Genome Initiative at Rutgers, Waksman Institute, Rutgers
 University
 190 Frelinghuysen Road, Piscataway, NJ 08854, USA
 Tel: 732 445 3801
 Fax: 732 445 5735
 Email: bhatti@waksman.rutgers.edu
 Seq primer: SP6
 Class: BAC ends
 High quality sequence start: 99.
 Location/Qualifiers
 1..380
 /organism="Zea mays"
 /mol_type="genomic DNA"
 /cultivar="B73"
 /db_xref="taxon:4577"
 /clone="ZMMBB0641H10"
 /lab_host="E. coli DH108"
 /clone_lib="ZMMBBB (HindIII)"
 /note="Vector: pCUGI; Site_1: HindIII; Site_2: HindIII"

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 Query Match 82.9%; Score 17.4; DB 9; Length 380;
 Best Local Similarity 94.7%; Pred. No. 1.1e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 CAGTGACATGCAGGCTCTAG 19
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 Db 210 CATTGACATGCAGGCTCTAG 192

RESULT 8
 CG176005 718 bp DNA linear GSS 21-AUG-2003
 LOCUS PUIWY95TD ZM 0.6 1.0_KB Zea mays genomic clone ZMMBtra0620021,
 DEFINITION genomic survey sequence.
 ACCESSION CG176005
 VERSION CG176005.1 GI:34066803
 KEYWORDS GSS.
 SOURCE Zea mays
 Zea mays
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.
 1 (bases 1 to 718)
 Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
 Renick,A., Frazer,C.M., Yuan,Y., San Miguel,P., Ma,J. and
 Bennetzen,J.
 Maize Genomics Consortium
 Unpublished (2003)
 Other GSSs: PUIWY95TB
 Contact: Cathy Whitelaw
 TIGR
 9712 Medical Center Drive, Rockville, MD 20850, USA
 Tel: 301-838-5843
 Fax: 301-838-0208

LOCUS BF607360 722 bp mRNA linear EST 01-APR-2001
 DEFINITION MY1.000302 Mouse 9-day fetus cdna library ICRPp522 Mus musculus
 cdna clone ICRPp522J2149 5', mRNA sequence.
 ACCESSION BF607360
 VERSION BF607360.1 GI:13503852
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 Yahyawi,M., Hennig,S., Neidhardt,L., Radelof,U., Hermann,B.G.,
 Lehrach,H. and O'Brien,J.
 Detection of a high number of novel genes in a 9-day mouse embryo
 cdna library normalised by oligonucleotide fingerprinting
 Unpublished (2001)
 Contact: Hennig S
 laboraty 123, dept.Lehrach
 Max-Planck-Institut fuer Molekulare Genetik
 Ihnestr.63-73, D-14195 Berlin, Germany
 Tel: +49 30 8413 1612
 Fax: +49 30 8413 1380
 Email: hennig@molgen.mpg.de
 EST's are made from clones being representatives of clone clusters.
 Clone clusters were calculated from oligonucleotide fingerprints.
 PCR Primers
 FORWARD: 5'-GAGCTATTCGAGTAGTAGTA-3'
 BACKWARD: 5'-TATACGACTCTACTATAGG-3'
 Seq primer: 5'-ATTAGGTGACATATAG-3'
 High quality sequence stop: 722.
 Location/Qualifiers
 1..722
 /organism="Mus musculus"
 /mol_type="mRNA"
 /db_xref="taxon:10090"
 /clone="ICRPp522J2149"
 /tissue_type="whole embryo"
 /dev_stage="embryonic 9-day"
 /lab_host="E.coli, XL1 blue"
 /clone_lib="Mouse 9-day fetus cdna library ICRPp522"
 /note="vector: PSVSPORT1; Site 1: NotI; Site 2: SalI;
 Library preparation by oligo dt priming of RNA. Clones can
 be ordered from the Resource Center in Berlin,
 http://www.rzpd.de."

ORIGIN
 Query Match 82.9%; Score 17.4; DB 2; Length 722;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Email: whitelaw@tigr.org
 Seq primer: TP
 Class: sheared ends.
 Location/Qualifiers
 1..718
 /organism="Zea mays"
 /mol_type="genomic DNA"
 /strain="B73"
 /db_xref="taxon:4577"
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 /clone_lib="ZM 0.6 1.0_KB"
 /note="Vector: PCR4-TOPO; Site_1: EcoRI; 0.6-1.0 kb high
 Cot selected genomic DNA library"

ORIGIN
 Query Match 82.9%; Score 17.4; DB 9; Length 718;
 Best Local Similarity 94.7%; Pred. No. 1.2e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 CAGTGACATGCAGGCTCTAG 19
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 Db 374 CATTGACATGCAGGCTCTAG 392

RESULT 9
 BF607360 722 bp mRNA linear EST 01-APR-2001
 LOCUS MY1.000302 Mouse 9-day fetus cdna library ICRPp522 Mus musculus
 DEFINITION cdna clone ICRPp522J2149 5', mRNA sequence.

ACCESSION BF607360
 VERSION BF607360.1 GI:13503852
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 Yahyawi,M., Hennig,S., Neidhardt,L., Radelof,U., Hermann,B.G.,
 Lehrach,H. and O'Brien,J.
 Detection of a high number of novel genes in a 9-day mouse embryo
 cdna library normalised by oligonucleotide fingerprinting
 Unpublished (2001)
 Contact: Hennig S
 laboraty 123, dept.Lehrach
 Max-Planck-Institut fuer Molekulare Genetik
 Ihnestr.63-73, D-14195 Berlin, Germany
 Tel: +49 30 8413 1612
 Fax: +49 30 8413 1380
 Email: hennig@molgen.mpg.de
 EST's are made from clones being representatives of clone clusters.
 Clone clusters were calculated from oligonucleotide fingerprints.
 PCR Primers
 FORWARD: 5'-GAGCTATTCGAGTAGTAGTA-3'
 BACKWARD: 5'-TATACGACTCTACTATAGG-3'
 Seq primer: 5'-ATTAGGTGACATATAG-3'
 High quality sequence stop: 722.
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 1..722
 /organism="Mus musculus"
 /mol_type="mRNA"
 /db_xref="taxon:10090"
 /clone="ICRPp522J2149"
 /tissue_type="whole embryo"
 /dev_stage="embryonic 9-day"
 /lab_host="E.coli, XL1 blue"
 /clone_lib="Mouse 9-day fetus cdna library ICRPp522"
 /note="vector: PSVSPORT1; Site 1: NotI; Site 2: SalI;
 Library preparation by oligo dt priming of RNA. Clones can
 be ordered from the Resource Center in Berlin,
 http://www.rzpd.de."

ORIGIN
 Query Match 82.9%; Score 17.4; DB 2; Length 722;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Best Local Similarity 90.0%; Pred. No. 1.2e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAGC 20
Db 314 CAGTANCATGCAGGTCTAGC 333

RESULT 10
CC340825 772 bp DNA linear GSS 16-MAY-2003
OGQAQ84TH ZM 0.7 1.5 KB Zea mays genomic clone ZMMBma0368M23,
genomic survey sequence.
ACCESSION CC340825
VERSION CC340825.1 GI:30810231
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
Consortium for Maize Genomics
Unpublished (2002)
Contact: Cathy Whitelaw
TIGR

TITLE
JOURNAL
COMMENT

FEATURES
source
1. .772
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone="ZMMBma0368M23"
/clone_lib="ZM 0.7 1.5 KB"
/notes="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN
Query Match 82.9%; Score 17.4; DB 8; Length 772;
Best Local Similarity 94.7%; Pred. No. 1.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAG 19
Db 744 CATTGACATGCAGGTCTAG 726

RESULT 11
CG211386 772 bp DNA linear GSS 22-AUG-2003
OGSCS03TC ZM 0.7 1.5 KB Zea mays genomic clone ZMMBma0832B05,
genomic survey sequence.
ACCESSION CG211386
VERSION CG211386.1 GI:34111216
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
Consortium for Maize Genomics
Unpublished (2002)
Contact: Cathy Whitelaw
TIGR

TITLE
JOURNAL
COMMENT

FEATURES
source
1. .772
/organism="Zea mays"
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/strain="B73"
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/notes="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN
Query Match 82.9%; Score 17.4; DB 8; Length 772;
Best Local Similarity 94.7%; Pred. No. 1.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAG 19
Db 744 CATTGACATGCAGGTCTAG 726

RESULT 12
CG362883 805 bp DNA linear GSS 26-AUG-2003
OGYBS24TH ZM 0.7 1.5 KB Zea mays genomic clone ZMMBma0643C23,
genomic survey sequence.
ACCESSION CG362883
VERSION CG362883.1 GI:34280150
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
Consortium for Maize Genomics
Unpublished (2002)
Other GSSs: OGYBS24TV
Contact: Cathy Whitelaw
TIGR

TITLE
JOURNAL
COMMENT

FEATURES
source
1. .805
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone="ZMMBma0643C23"
/clone_lib="ZM 0.7 1.5 KB"
/notes="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN
Query Match 82.9%; Score 17.4; DB 9; Length 805;
Best Local Similarity 94.7%; Pred. No. 1.2e+03;

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```

TITLE
JOURNAL
COMMENT
Consortium for Maize Genomics
Unpublished (2002)
Contact: Cathy Whitelaw
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TP
Class: sheared ends.
Location/Qualifiers
1. .772
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
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/clone_lib="ZM 0.7 1.5 KB"
/notes="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN
Query Match 82.9%; Score 17.4; DB 9; Length 772;
Best Local Similarity 94.7%; Pred. No. 1.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCTAG 19
Db 287 CATTGACATGCAGGTCTAG 305

RESULT 12
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LOCUS
DEFINITION OGYBS24TH ZM 0.7 1.5 KB Zea mays genomic clone ZMMBma0643C23,
genomic survey sequence.
ACCESSION CG362883
VERSION CG362883.1 GI:34280150
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
Consortium for Maize Genomics
Unpublished (2002)
Other GSSs: OGYBS24TV
Contact: Cathy Whitelaw
TIGR

TITLE
JOURNAL
COMMENT

FEATURES
source
1. .805
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
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/clone_lib="ZM 0.7 1.5 KB"
/notes="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN
Query Match 82.9%; Score 17.4; DB 9; Length 805;
Best Local Similarity 94.7%; Pred. No. 1.2e+03;

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Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 CAGTGACATGCAGGCTCTAG 19
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Db 384 CATTGACATGCAGGCTCTAG 366

RESULT 13
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LOCUS
DEFINITION 603189492F1 NIH_MGC_95 Homo sapiens cDNA clone IMAGE:5260847 5',
            mRNA sequence.
ACCESSION BI548151
VERSION BI548151.1 GI:15435463
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 816)
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
          Email: cgsabbs-remail.nih.gov
          Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
          cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
          Tohiyuki and Piero Carninci (RIKEN)
          cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
          DNA Sequencing by: Incyte Genomics, Inc.
          Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          http://image.llnl.gov
          Plate: LLAM11657 row: g column: 24
          High quality sequence stop: 740.

FEATURES
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                /clone="IMAGE:5260847"
                /tissue_type="hippocampus"
                /lab_host="DH10B"
                /clone_libs="NIH MGC 95"
                /note="Organ: brain; Vector: pBluescriptR (modified
                pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI
                (gtcgag); Oligo-dT primed using primer
                5'-TTTTTTTTTTTTTTVN-3', size-selected for average
                insert size 2.5 kb and normalized to ROT 5. This is a
                primary library enriched for full-length clones and
                constructed using the Cap-trapper method (Carninci, in
                preparation). Library constructed by M. Brownstein
                (NIH/NHGRI, National Institutes of Health). Note: this
                is a NIH_MGC Library."

ORIGIN
    Query Match 82.9%; Score 17.4; DB 4; Length 816;
    Best Local Similarity 94.7%; Pred. No. 1.2e+03;
    Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 CAGTGACATGCAGGCTCTAG 19
    |||||||
Db 295 CAGTGACATGCAGGCTCTAG 277

RESULT 14
CC340835
LOCUS
DEFINITION CC340835 OGAQA84TV ZM_0.7_1.5 KB Zea mays genomic clone ZMMBMA0368M23,
            genomic survey sequence.
ACCESSION CC340835
VERSION CC340835.1 GI:30810241
KEYWORDS GSS.

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 CAGTGACATGCAGGCTCTAG 19
    |||||||
Db 295 CAGTGACATGCAGGCTCTAG 277

RESULT 14
CC340835
LOCUS
DEFINITION CC340835 OGAQA84TV ZM_0.7_1.5 KB Zea mays genomic clone ZMMBMA0368M23,
            genomic survey sequence.
ACCESSION CC340835
VERSION CC340835.1 GI:30810241
KEYWORDS GSS.

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SOURCE ORGANISM
Zea mays
Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 817)
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
          Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
          Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
TITLE Consortium for Maize Genomics
JOURNAL Unpublished (2002)
COMMENT Contact: Cathy Whitelaw
          TIGR
          9712 Medical Center Drive, Rockville, MD 20850, USA
          Tel: 301-838-5843
          Fax: 301-838-0208
          Email: whitelaw@tigr.org
          Seq primer: TF
          Class: sheared ends.
          Location/Qualifiers
              1..817
                  /organism="Zea mays"
                  /mol_type="genomic DNA"
                  /strain="B73"
                  /db_xref="taxon:4577"
                  /clone="ZMMBMA0368M23"
                  /clone_lib="ZM 0.7_1.5 KB"
                  /note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
                  methylation filtered genomic DNA library"

ORIGIN
    Query Match 82.9%; Score 17.4; DB 8; Length 817;
    Best Local Similarity 94.7%; Pred. No. 1.2e+03;
    Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 CAGTGACATGCAGGCTCTAG 19
    |||||||
Db 488 CATTGACATGCAGGCTCTAG 506

RESULT 15
CG211861/c
LOCUS
DEFINITION CG211861 OGIAM29TV ZM_0.7_1.5 KB Zea mays genomic clone ZMMBMA0720F09,
            genomic survey sequence.
ACCESSION CG211861
VERSION CG211861.1 GI:34111691
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 923)
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
          Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
          Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
TITLE Consortium for Maize Genomics
JOURNAL Unpublished (2002)
COMMENT Other GSSs: OGIAM29TH
          Contact: Cathy Whitelaw
          TIGR
          9712 Medical Center Drive, Rockville, MD 20850, USA
          Tel: 301-838-5843
          Fax: 301-838-0208
          Email: whitelaw@tigr.org
          Seq primer: TF
          Class: sheared ends.
          Location/Qualifiers
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                  /mol_type="genomic DNA"
                  /strain="B73"

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methylation filtered genomic DNA library"

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Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTCACATGCAGGTCCTAG 19
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Db 157 CATTGCACATGCAGGTCCTAG 139

RESULT 16
BY609340
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

Mus musculus
Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 382)
Okazaki, Y., Furuno, M., Saito, R., Suzuki, H., Yamanaka, I.,
Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A.,
Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D.P., Bult, C.,
Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuoka, H.,
Batalov, S., Beisel, K.W., Blake, J.A., Bradt, D., Brusic, V.,
Chothia, C., Corbani, L.E., Cousins, S., Dalla, E., Dragani, T.A.,
Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S.,
Kawaji, H., Kwasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A.,
Kurochkin, I.V., Lee, Y., Lenhard, B., Lyons, P.A., Maglott, D.R.,
Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T.,
Numata, K., Okido, T., Pavan, W.J., Pertea, G., Pesole, G.,
Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramachandran, S.,
Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ringwald, M.,
Sandelin, A., Schneider, C., Sempile, C.A., Setou, M., Shimada, K.,
Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M.,
Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y.,
Wells, C., Wilming, L.G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I.,
Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P.,
Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M.,
Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
Arakawa, T., Fukuda, S., Harai, A., Hashizume, W., Imotani, K., Ishii, Y.,
Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K.,
Shinagawa, A., Yasuniishi, A., Yoshino, M., Waterston, R., Landet, E.S.,
Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
1245683
12456851
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
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The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
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Fax: 81-45-503-9216
Email: genome-res@gsc.riken.jp, URL: http://genome.gsc.riken.jp/
Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S.,
Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H.,
Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R.,
Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K.,

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Shiraki, T., Tagami, M., Waki, K., Watahiki, A., Muramatsu, M. and
Hayashizaki, Y. Direct Submission
Computational Analysis of Full-Length Mouse cDNAs Compared with
Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
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encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Tissues were provided by Michela Fagioli and Takao K. Hensch (
Laboratory for Neuronal Circuit Development Brain Science Institute
RIKEN 2-1 Hirosawa Wako-shi, Saitama 351-0198 Japan ) whose
assistance we gratefully acknowledge.
Please visit our web site (http://genome.gsc.riken.go.jp) for
further details.
FEATURES
Location/Qualifiers
source
1..382
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="K230308M20"
/tissue_type="visual cortex"
/clone_lib="RIKEN full-length enriched, visual cortex"

ORIGIN
Query Match      81.0%; Score 17; DB 6; Length 382;
Best Local Similarity 100.0%; Pred. No. 1.7e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 AGTCACATGCAGGTCCTA 18
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Db 28 AGTCACATGCAGGTCCTA 44

RESULT 17
BY649695
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 408)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Osato, N., Saito, R., Suzuki, H., Yamanaka, I.,
Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A.,
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Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuoka, H.,
Batalov, S., Beisel, K.W., Blake, J.A., Bradt, D., Brusic, V.,
Chothia, C., Corbani, L.E., Cousins, S., Dalla, E., Dragani, T.A.,
Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T.,
Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S.,
Kawaji, H., Kwasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A.,
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Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramachandran, S.,
Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ringwald, M.,

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Sandelin,A., Schneider,C., Semple,C.A., Setou,M., Shimada,K., Sultana,R., Takenaka,Y., Taylor,M.S., Teasdale,R.D., Tomita,M., Verardo,R., Wagner,L., Wallestedt,C., Wang,Y., Watanabe,Y., Wells,C., Wilming,L.G., Wynshaw-Boris,A., Yanagisawa,M., Yang,I., Yang,L., Yuan,Z., Zavolan,M., Zhu,Y., Zimmer,A., Carninci,P., Hayatsu,N., Hirozane-Kishikawa,T., Konno,H., Nakamura,M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Aizawa,K., Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imotani,K., Ishii,Y., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata,K., Shinaawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander,E.S., Rogers,J., Birney,E. and Hayashizaki,Y.

Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs

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12466851

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The Institute of Physical and Chemical Research (RIKEN)

1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan

Tel: 81-45-503-9222

Fax: 81-45-503-9216

Email: genome-res@gsc.riken.jp, URL:http://genome.gsc.riken.jp/

Aizawa,K., Akimura,T., Arakawa,T., Carninci,P., Fukuda,S., Hirozane,T., Imotani,K., Ishii,Y., Itoh,M., Kawai,J., Konno,H., Miyazaki,A., Murata,M., Nakamura,M., Nomura,K., Numazaki,R., Ohno,M., Sakai,K., Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shiraki,T., Tagami,M., Waki,K., Watahiki,A., Muramatsu,M. and Hayashizaki,Y. Direct Submission

TITLE

JOURNAL
MEDLINE
PUBMED
COMMENT

Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

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Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

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Tissues were provided by Michela Fagiolini and Takao K. Hensch (Laboratory for Neuronal Circuit Development Brain Science Institute RIKEN 2-1 Hirotsawa,Wako-shi,Saitama 351-0198 Japan) whose assistance we gratefully acknowledge.

Please visit our web site (http://genome.gsc.riken.go.jp) for further details.

FEATURES

source
1. .408
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="K330043P16"
/tissue_type="visual cortex"
/clone_lib="RIKEN full-length enriched, visual cortex"

ORIGIN

Query Match 81.0%; Score 17; DB 6; Length 408;
Best Local Similarity 100.0%; Pred. No. 1.7e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 AGTGACATGCAGGCTTA 18
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Db 56 AGTGACATGCAGGCTTA 72
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RESULT 18

CA124343/c

LOCUS

DEFINITION

SCQGLR1086F09.9 LRI Saccharum officinarum cDNA clone SCQGLR1086F09

5', mRNA sequence.

ACCESSION

CA124343

VERSION

CA124343.1 GI:34977651

KEYWORDS

Saccharum officinarum

SOURCE

EST.

ORGANISM

Saccharum officinarum

REFERENCE

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location/Qualifiers

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location/Qualifiers

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location/Qualifiers

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Akinret, B., Levins, M., McGann, S., Tsegaye, G., Geer, K., Krol, M., de Jong, P., and Fraser, C.M.
 Mouse BAC End Sequences from Library RPCI-23
 Unpublished (1999)
 Other_GSSs: RPCI-23-435G22.TV
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org

Clones are derived from the mouse BAC library RPCI-23. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/orderingframe.htm>) or from Resea ch Genetics (info@resgen.com). BAC end page: http://www.tigr.org/tdb/bac_ends/mouse/bac_end_intro.html
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 Seq primer: SP6
 Class: BAC ends.

FEATURES

Location/Qualifiers
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 /sex="Female"
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/note="Organ: Kidney/Brain; Vector: pBACe3.6; Site 1: EcoRI; Site 2: EcoRI; Female C57BL/6J mouse kidney and/or brain genomic DNA was isolated and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBACe3.6 vector at the EcoRI sites. The ligation products were transformed into DH10B electrocompetent cells (BRL Life Technologies)."

ORIGIN

Query Match 81.0%; Score 17; DB 8; Length 633;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCT 17
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Db 122 CAGTGACATGCAGGTCT 138

RESULT 20

BB650662/c

LOCUS

DEFINITION BB650662 RIKEN full-length enriched, 0 day neonate cerebellum Mus musculus cDNA clone C230020D16 5', mRNA sequence.

ACCESSION BB650662

VERSION BB650662.1

KEYWORDS GI:16484917

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 642)

REFERENCE

AUTHORS

Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T., Konno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K., Ohno, H., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F., Takeda, Y., Tanaka, T., Toya, T., Muramatsu, M. and Hayashizaki, Y.
 RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
 Unpublished (2001)

TITLE

JOURNAL

COMMENT

Contact: Yoshihide Hayashizaki

Laboratory for Genome Exploration Research Group, RIKEN Genomic

Sciences Center(GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel: 81-45-503-9222
 Fax: 81-45-503-9216

Email: genome-res@gsc.riken.jp, URL:<http://genome.gsc.riken.jp/>
 Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
 Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
 RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y.

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
 Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamanaka, I., Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and Hayashizaki, Y.

Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences. Mamm. Genome. 12, 673-677 (2001)
 Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

e mouse tissues.

FEATURES

Location/Qualifiers
 1..642
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 /lab_host="DH10B"
 /clone_lib="RIKEN full-length enriched, 0 day neonate cerebellum"

/note="Site 1: Sali; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5'
 GAGAGAGAGAGATCCAGAGCTCTTTTTTTTTTTTNN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 20.0 and subtraction to Rot = 479.0. Second strand cDNA was prepared with the primer adaptor of sequence [5' GAGAGAGAGATTCGAGTTAATAATATCCCTCCCCC 3']. cDNA was cleaved with XhoI and BamHI. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I."

ORIGIN

Query Match 81.0%; Score 17; DB 2; Length 642;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGTCT 17
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Db 27 CAGTGACATGCAGGTCT 11

RESULT 21

BB293162/c

LOCUS

655 bp mRNA linear EST 24-OCT-2001

DEFINITION BB293162 RIKEN full-length enriched, 9.5 days embryo parthenogenote
Mus musculus cDNA clone B130017A20 3' similar to AK001617 Homo
sapiens cDNA FLJ10755 fis, clone NT2RP3004569, weakly similar to
ANKYRIN, BRAIN VARIANT 1, mRNA sequence.

ACCESSION BB293162
VERSION BB293162.2 GI:16401650
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T.,
Hara, A., Hiramoto, K., Hori, F., Ichi, Y., Ito, M., Kawai, J.,
Konno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K.,
Ohno, H., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K.,
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,
Takeda, Y., Tanaka, T., Toyota, T., Muramatsu, M. and Hayashizaki, Y.
RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
Unpublished (2001)
On Jul 10, 2000 this sequence version replaced gi:8993654.
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gscc.riken.jp, URL: http://genome.gsc.riken.jp/
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
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Normalization and subtraction of cap-trapper-selected cDNAs to
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wagi, K., Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
Watahiki, M., Yoneda, Y., Iihikawa, T., Ozawa, K., Tanaka, T.,
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Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamanaka, I.,
Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and
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Please visit our web site (http://genome.gsc.riken.go.jp/) for
further details.

FEATURES
source
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/organism="Mus musculus"
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/clone_lib="RIKEN full-length enriched, 9.5 days embryo
parthenogenote"
/note="Site 1: Sali; Site 2: BamHI; cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken

Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
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GAGAGAGAGAGATCCAGAGCTCTTTTTTTTTTTTNN 3'], cDNA was
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. cDNA went through one round of subtraction to
Rot = 229.0 Second strand cDNA was prepared with the
primer adapter of sequence [5',
GAGAGAGAGATTCGAGTTAATTAATCCCTCCCCCCCC 3']. cDNA
was cleaved with XhoI and BamHI. Vector: a modified
pBluescript KS(+) after bulk excision from Lambda PLC I."

ORIGIN
Query Match 81.0%; Score 17; DB 2; Length 655;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CAGTGACATGCAGGTCT 17
|||||||
Db 145 CAGTGACATGCAGGTCT 129

RESULT 22
AG457137
LOCUS AG457137.1 GI:48148651
DEFINITION Mus musculus molossinus DNA, clone:MSMg01-343P07.TJ, genomic survey
sequence.
ACCESSION AG457137
VERSION AG457137.1
KEYWORDS GSS.
SOURCE Mus musculus molossinus
ORGANISM Mus musculus molossinus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1
Hattori, M., Toyoda, A., Noguchi, H., Kojima, T. and Sakaki, Y.
BAC end Sequences of Library MSMg01
Unpublished
2 (bases 1 to 768)
Direct Submission
Submitted (17-NOV-2003) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail: hattori@gscc.riken.jp, URL: http://hgpc.riken.go.jp/
Tel: 81-45-503-9111, Fax: 81-45-503-9170)
Clones are derived from the mouse BAC library MSMg01. For BAC
library availability, please contact Kuniya Abe (abe@rtc.riken.jp).
The Institute of Physical and Chemical Research (RIKEN) 3-1-1
Koyadai, Tsukuba, 305-0074 Japan
phone: 81-298-36-9189, fax: 81-298-36-9199
e-mail: abe@rtc.riken.jp

PRIMERS
LIBRARY Sequencing : TJ
Vector : pBACe3.6
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R.Site 2 : EcoRI
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TITLE Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE 20499374
PUBMED 11042159
REFERENCE

AUTHORS Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P., Konno, H., Akiyama, J., Nishi, K., Kitsuai, T., Tashiro, H., Itoh, M., Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A., Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watabiki, M., Yoneda, Y., Ishikawa, T., Osawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
TITLE RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multipicillary sequencer
JOURNAL Genome Res. 10 (11), 1757-1771 (2000)
MEDLINE 20530913
PUBMED 11076861
REFERENCE

AUTHORS 4 The RIKEN Genome Exploration Research Group Phase II Team and the FANTOM Consortium.
TITLE Functional annotation of a full-length mouse cDNA collection
JOURNAL Nature 409, 685-690 (2001)
REFERENCE 5

AUTHORS The FANTOM Consortium and the RIKEN Genome Exploration Research Group Phase I & II Team.
TITLE Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs
JOURNAL Nature 420, 563-573 (2002)
REFERENCE 6 (bases 1 to 2445)

AUTHORS Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W., Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T., Katoh, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M., Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y.
TITLE Direct Submission

JOURNAL Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.jp, URL: <http://genome.gsc.riken.jp/>, Tel: 81-45-503-9222, Fax: 81-45-503-9216)

COMMENT cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Tissues were provided by Dr. Tomohiro Kono (Department of Animal Science, Tokyo University of Agriculture, 1737 Hunko Atsugi City, Kanagawa Prefecture, Japan) whose assistance we gratefully acknowledge.

Please visit our web site for further details.

URL: <http://genome.gsc.riken.jp/>

URL: <http://fantom.gsc.riken.jp/>

Location/Qualifiers

1. 2445

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ORIGIN

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Qy 1 CAGTGACATGCAGGTCT 17

|||||

Db 1935 CAGTGACATGCAGGTCT 1919

Search completed: September 6, 2005, 21:56:00
 Job time : 1506.84 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 735.656 Seconds
(without alignments)
1383.200 Million cell updates/sec

Title: US-10-729-421-40
Perfect score: 21
Sequence: 1 cagtgcacatgcaggtctagct 21

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database :

GenEmbl.*

1: gb_ba.*

2: gb_htg.*

3: gb_in.*

4: gb_on.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pt.*

10: gb_ro.*

11: gb_sta.*

12: gb_by.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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| C 5 | 18.4 | 87.6 | 161516 | 2 | CR391906 Danio rer |
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| C 29 | 17.4 | 82.9 | 110000 | 2 | Continuation (2 of |
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ALIGNMENTS
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ACCESSION AC109544
VERSION AC109544.5 GI:25006749
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SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 232064)
REFERENCE
AUTHORS Murzyn,D,Marie., Metzker,M,Lee., Abramzon,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
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Claveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
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Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wleczyk,R., Wooden,J., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,

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TITLE
JOURNAL
REFERENCE
AUTHORS
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AUTHORS
TITLE
JOURNAL
COMMENT

Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von
Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O.,
Weinstock, G. and Gibbs, R. A.
Direct Submission
Unpublished
2 (bases 1 to 232064)
Worley, K. C.
Direct Submission
Submitted (05-FEB-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 232064)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (15-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Nov 15, 2002 this sequence version replaced gi:23266105.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: QGBR
Center clone name: CH230-202010
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 210440 bases at least Q40
Consensus quality: 213054 bases at least Q30
Consensus quality: 214694 bases at least Q20
Estimated insert size: 221175; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation
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* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
*
* 1 232064: contig of 232064 bp in length.
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ORIGIN

Query Match 92.4%; Score 19.4; DB 2; Length 232064;
Best Local Similarity 95.2%; Pred. NO. 63;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTTAGCT 21

Db 34510 CAGTGACATGCAGGCTTAGCT 34530

RESULT 2

AC133226/c
LOCUS AC133226 258823 bp DNA linear HTG 15-NOV-2002
DEFINITION Rattus norvegicus clone CH230-329C22, *** SEQUENCING IN PROGRESS

ACCESSION

AC133226 GI:25007420

VERSION HTG; HTGS PHASE2; HTGS_DRAFT; HTGS_ENRICHED.

KEYWORDS Rattus norvegicus (Norway rat)

SOURCE

ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 258823)

REFERENCE

1 Muzny, D. Waré, Metzker, M. Lee., Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Albrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Ayagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Blawie, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Cesar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T. T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S. L., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idiebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpachy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhow, L., Loulseghe, H., Lozada, R. J., Lu, X., Ma, J., Maheshwari, M., Mahindartine, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhney, S., McLeod, M. P., McNeill, T. Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankovis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwakoleme, O., Okwionu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Pappert, F., Poindexter, A., Popovic, D., Primus, E., Pu, L., Pu, L., Puazo, M., Quiroz, J., Rachin, E., Reeves, K., Regier, M. A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, P., Rivers, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S. J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sieson, I., Sitter, C. D., Smajls, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villalana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,

Williams, G., Willson, R., Wlezyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, X., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O., Weinstock, G., and Gibbs, R. A.

Direct Submission

Unpublished

2 (bases 1 to 258823)

Rat Genome Sequencing Consortium.

Direct Submission

Submitted (08-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 258823)

Rat Genome Sequencing Consortium.

Direct Submission

Submitted (15-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

COMMENT

On Nov 15, 2002 this sequence version replaced gi:22771302. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center of Medicine

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: KBNW

Center clone name: CH230-329C22

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 187552 bases at least Q40

Consensus quality: 190690 bases at least Q30

Consensus quality: 192289 bases at least Q20

Estimated insert size: 192259; sum-of-contigs estimation

Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 1 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces

* is believed to be correct as given, however the sizes

* of the gaps between them are based on estimates that have

* provided by the submitter.

* This sequence will be replaced

* by the finished sequence as soon as it is available and

* the accession number will be preserved.

* the accession number will be 258823 bp in length.

FEATURES
source

1. 258823

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clones="CH230-329C22"

1. 1057

/note="wgs end extension

clone_end:Sp6"

5970_6869

/note="clone boundary

clone_end:Sp6

site:

misc_feature

misc_feature


```

VERSION      AP000772.2  GI:8118931
KEYWORDS     HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE       Homo sapiens (human)
ORGANISM     Homo sapiens
              Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE    1 (bases 1 to 155858)
AUTHORS      Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
              Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
              Published Only in Database (1999)
TITLE        2 (bases 1 to 155858)
JOURNAL      Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
              Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
              Direct Submission
AUTHORS      Submitted (25-NOV-1999) Masahira Hattori, The Institute of Physical
              and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
              Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
              Japan (E-mail:hattori@gsc.riken.go.jp,
              URL:http://hgp.gsc.riken.go.jp/, Tel:81-42-778-9923,
              Fax:81-42-778-9924)
COMMENT      On May 31, 2000 this sequence version replaced gi:6997610.
              ----- Genome Center
              Center: RIKEN Genomic Sciences Center (GSC)
              Center code: RIKEN
              Web site: http://hgp.gsc.riken.go.jp/
              Contact: hattori@gsc.riken.go.jp
              ----- Project Information
              Center project name: HumDraft11
              Center Clone name: CMB9-7B14
              ----- Summary Statistics
              Sequencing vector: PCR products; 100% of reads
              Chemistry: Dye-terminator ET-amersham; 100% of reads
              Assembly program: Phrap; version 0.990329
              Consensus quality: 135660 bases at least Q40
              Consensus quality: 145354 bases at least Q30
              Consensus quality: 150658 bases at least Q20
              Insert size: 153258; sum-of-Contigs
              Quality coverage: 4.31x in Q20 bases; sum-of-contigs
              -----
NOTE: This is a 'working draft' sequence. It currently consists of
27 contigs. The true order of the pieces is not known and their
order in this sequence record is arbitrary. Gaps between the
contigs are represented as runs N, but the exact sizes of the gaps
are unknown. This record will be updated with the finished sequence
as soon as it is available and the accession number will be
preserved
1
18489 contig of 18489 bp in length
18590 contig of 17553 bp in length
36243 contig of 13655 bp in length
49998 contig of 13726 bp in length
63824 contig of 12968 bp in length
76892 contig of 9036 bp in length
86028 contig of 6745 bp in length
93157 contig of 7029 bp in length
99902 contig of 6745 bp in length
100001 contig of 100 bp
100002 contig of 6584 bp in length
106586 contig of 100 bp
106586 contig of 100 bp
111187 contig of 4502 bp in length
111188 contig of 100 bp
111287 contig of 100 bp
112288 contig of 3919 bp in length
115207 contig of 100 bp
115307 contig of 4571 bp in length
119878 contig of 100 bp
119977 contig of 100 bp
124080 contig of 4103 bp in length
124081 contig of 100 bp
124181 contig of 100 bp
127016 contig of 2835 bp in length
127116 contig of 100 bp
129487 contig of 2372 bp in length
129488 contig of 100 bp
129587 contig of 3436 bp in length
13023 contig of 3436 bp in length
133024 contig of 100 bp
133123 contig of 3209 bp in length
133124 contig of 100 bp
136332 contig of 100 bp
136432 contig of 100 bp
136684 contig of 3252 bp in length
139685 contig of 100 bp
139785 contig of 3132 bp in length
142916 contig of 3132 bp in length
143016 contig of 100 bp
143017 contig of 2153 bp in length
145169 contig of 2153 bp in length
145170 contig of 100 bp
145270 contig of 100 bp
147638 contig of 2369 bp in length
147738 contig of 100 bp
147739 contig of 1165 bp in length
148904 contig of 100 bp
149003 contig of 100 bp
149004 contig of 1378 bp in length
150381 contig of 100 bp
150482 contig of 1196 bp in length
151677 contig of 1196 bp in length
151678 contig of 100 bp
151778 contig of 1810 bp in length
153588 contig of 100 bp
153688 contig of 1011 bp in length
154699 contig of 100 bp
154799 contig of 1060 bp in length.
155858 contig of 1060 bp in length.
FEATURES             Location/Qualifiers
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                        /organism="Homo sapiens"
                        /mol_type="genomic DNA"
                        /db_xref="taxon:9606"
                        /chromosome="11"
                        /map="11q22"
     misc_feature      1..18489
                        /note="assembly_fragment"

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misc_feature 18590..36142
/note="assembly_fragment"
misc_feature 36243..49897
/note="assembly_fragment"
misc_feature 49998..63723
/note="assembly_fragment"
misc_feature 63824..76791
/note="assembly_fragment"
misc_feature 76892..85927
/note="assembly_fragment"
misc_feature 86028..93056
/note="assembly_fragment"
misc_feature 93157..99901
/note="assembly_fragment"
misc_feature 100002..106585
/note="assembly_fragment"
misc_feature 106686..111187
/note="assembly_fragment"
misc_feature 111288..115206
/note="assembly_fragment"
misc_feature 115307..119877
/note="assembly_fragment"
misc_feature 119978..124080
/note="assembly_fragment"
misc_feature 124181..127015
/note="assembly_fragment"
misc_feature 127116..129487
/note="assembly_fragment"
misc_feature 129588..133023
/note="assembly_fragment clone_end:SP6 vector_side:left"
misc_feature 133124..136332
/note="assembly_fragment"
misc_feature 136433..139684
/note="assembly_fragment"
misc_feature 139785..142916
/note="assembly_fragment"
misc_feature 143017..145169
/note="assembly_fragment"
misc_feature 145270..147638
/note="assembly_fragment"
misc_feature 147739..148903
/note="assembly_fragment"
misc_feature 149004..150381
/note="assembly_fragment"
misc_feature 150482..151677
/note="assembly_fragment"
misc_feature 151778..153587
/note="assembly_fragment"
misc_feature 153688..154698
/note="assembly_fragment"

Query Match 87.6%; Score 18.4; DB 2; Length 155858;
Best Local Similarity 95.0%; Pred.No. 2.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTCACATGCAGGCTAGC 20
DB 89083 CAGTCACATGCAGGCTAGC 89064

RESULT 5
CR391906/c
LOCUS CR391906 161516 bp DNA linear HTG 24-APR-2004
DEFINITION Danio rerio clone DKEY-211K10, *** SEQUENCING IN PROGRESS ***, 9
unordered pieces.
ACCESSION CR391906
VERSION CR391906.2 GI:46559615
KEYWORDS HTG; HTGS PHASE1
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.

```

```

REFERENCE 1 (bases 1 to 161516)
AUTHORS McLay, K.
JOURNAL Direct Submission
TITLE Submitted (23-APR-2004) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
COMMENT On Apr 24, 2004 this sequence version replaced gi:46517964.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: zfish-help@sanger.ac.uk
----- Project Information
Center project name: zK211K10
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: Dye-terminator; 100% of reads
Consensus quality: 159202 bases at least Q40
Consensus quality: 159648 bases at least Q30
Consensus quality: 160053 bases at least Q20
Insert size: 160716; sum-of-contigs
Insert size: 167493; 3.3% error; agarose-fp
Quality coverage: 6.27x in Q20 bases; sum-of-contigs Quality
coverage: 6.01x in Q20 bases; agarose-fp
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 9 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 2483: contig of 2483 bp in length
* 2484 2583: gap of 100 bp
* 2584 38899: contig of 36316 bp in length
* 38900 38999: gap of 100 bp
* 39000 55041: contig of 16042 bp in length
* 55042 55141: gap of 100 bp
* 55142 58203: contig of 3062 bp in length
* 58204 58304: gap of 100 bp
* 58304 70412: contig of 12109 bp in length
* 70413 70512: gap of 100 bp
* 70513 100144: contig of 29632 bp in length
* 100145 100244: gap of 100 bp
* 100245 110317: contig of 10073 bp in length
* 110318 110417: gap of 100 bp
* 110418 122844: contig of 12427 bp in length
* 122845 122944: gap of 100 bp
* 122945 161516: contig of 38572 bp in length.
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FEATURES
source
1..161516
/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clone="DKEY-211K10"
/clone_lib="DanioKey"
1..2483
/note="assembly fragment:00070
fragment_chain:1"
2584..38899
/note="assembly fragment:01225
fragment_chain:1"
39000..55041
/note="assembly fragment:00464
fragment_chain:1"
55142..58203
/note="assembly fragment:00087
fragment_chain:1"
58304..70412
/note="assembly fragment:00114
fragment_chain:2"
70513..100144
misc_feature
misc_feature
misc_feature
misc_feature
misc_feature
misc_feature

```


* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.

* 1 949: contig of 949 bp in length
* 950 1049: gap of 100 bp
* 1050 2006: contig of 957 bp in length
* 2007 2106: gap of 100 bp
* 2107 3119: contig of 1013 bp in length
* 3120 3219: gap of 100 bp
* 3220 4172: contig of 953 bp in length
* 4173 4272: gap of 100 bp
* 4273 5244: contig of 972 bp in length
* 5245 5344: gap of 100 bp
* 5345 6288: contig of 944 bp in length
* 6289 6388: gap of 100 bp
* 6389 7363: contig of 975 bp in length
* 7364 7463: gap of 100 bp
* 7464 8447: contig of 984 bp in length
* 8448 8547: gap of 100 bp
* 8548 9561: contig of 1014 bp in length
* 9562 9661: gap of 100 bp
* 9662 10597: contig of 936 bp in length
* 10598 10697: gap of 100 bp
* 10698 11718: contig of 1021 bp in length
* 11719 11818: gap of 100 bp
* 11819 12831: contig of 1013 bp in length
* 12832 12931: gap of 100 bp
* 12932 13900: contig of 989 bp in length
* 13901 14000: gap of 100 bp
* 14001 14953: contig of 953 bp in length
* 14954 15053: gap of 100 bp
* 15054 16082: contig of 1029 bp in length
* 16083 16182: gap of 100 bp
* 16183 17195: contig of 1013 bp in length
* 17196 17295: gap of 100 bp
* 17296 18344: contig of 1049 bp in length
* 18345 18444: gap of 100 bp
* 18445 19385: contig of 941 bp in length
* 19386 19485: gap of 100 bp
* 19486 20455: contig of 970 bp in length
* 20456 20555: gap of 100 bp
* 20556 21548: contig of 993 bp in length
* 21549 21648: gap of 100 bp
* 21649 22631: contig of 983 bp in length
* 22632 22731: gap of 100 bp
* 22732 23705: contig of 974 bp in length
* 23706 23805: gap of 100 bp
* 23806 24762: contig of 957 bp in length
* 24763 24862: gap of 100 bp
* 24863 25835: contig of 973 bp in length
* 25836 25935: gap of 100 bp
* 25936 26918: contig of 983 bp in length
* 26919 27018: gap of 100 bp
* 27019 28003: contig of 985 bp in length
* 28004 28103: gap of 100 bp
* 28104 29109: contig of 1006 bp in length
* 29110 29209: gap of 100 bp
* 29210 30234: contig of 1025 bp in length
* 30235 30334: gap of 100 bp
* 30335 31327: contig of 993 bp in length
* 31328 31427: gap of 100 bp
* 31428 32398: contig of 971 bp in length
* 32399 32498: gap of 100 bp
* 32499 33501: contig of 1003 bp in length
* 33502 33601: gap of 100 bp
* 33602 34612: contig of 1011 bp in length
* 34613 34712: gap of 100 bp
* 34713 35718: contig of 1006 bp in length
* 35719 35818: gap of 100 bp
* 35819 36820: contig of 1002 bp in length
* 36821 36920: gap of 100 bp

* 36921 37914: contig of 994 bp in length
* 37915 38014: gap of 100 bp
* 38015 38989: contig of 975 bp in length
* 38990 39089: gap of 100 bp
* 39090 40045: contig of 956 bp in length
* 40046 40145: gap of 100 bp
* 40146 41107: contig of 982 bp in length
* 41108 42108: gap of 100 bp
* 42109 42219: contig of 1012 bp in length
* 42220 42319: gap of 100 bp
* 42320 43325: contig of 1006 bp in length
* 43326 43425: gap of 100 bp
* 43426 44439: contig of 1014 bp in length
* 44440 44539: gap of 100 bp
* 44540 45494: contig of 955 bp in length
* 45495 45594: gap of 100 bp
* 45595 46580: contig of 986 bp in length
* 46581 46680: gap of 100 bp
* 46681 47695: contig of 1015 bp in length
* 47696 47795: gap of 100 bp
* 47796 48789: contig of 994 bp in length
* 48790 48890: gap of 100 bp
* 48891 49831: contig of 942 bp in length
* 49832 49932: gap of 100 bp
* 49933 50926: contig of 995 bp in length
* 50927 51026: gap of 100 bp
* 51027 52016: contig of 990 bp in length
* 52017 52116: gap of 100 bp
* 52117 53074: contig of 958 bp in length
* 53075 53174: gap of 100 bp
* 53175 54218: contig of 1044 bp in length
* 54219 54318: gap of 100 bp
* 54319 55335: contig of 1017 bp in length
* 55336 56368: contig of 933 bp in length
* 56369 57441: contig of 973 bp in length
* 57442 57541: gap of 100 bp
* 57542 58519: contig of 978 bp in length
* 58520 58619: gap of 100 bp
* 58620 59585: contig of 966 bp in length
* 59586 60652: contig of 967 bp in length
* 60653 60752: gap of 100 bp
* 60753 61736: contig of 984 bp in length
* 61737 62768: contig of 932 bp in length
* 62769 62868: gap of 100 bp
* 62869 63880: contig of 1012 bp in length
* 63881 63980: gap of 100 bp
* 63981 64974: contig of 994 bp in length
* 64975 65074: gap of 100 bp
* 65075 66085: contig of 1011 bp in length
* 66086 66185: gap of 100 bp
* 66186 67121: contig of 936 bp in length
* 67122 67221: gap of 100 bp
* 67222 68235: contig of 1014 bp in length
* 68236 68335: gap of 100 bp
* 68336 69321: contig of 986 bp in length
* 69322 69421: gap of 100 bp
* 69422 70456: contig of 1035 bp in length
* 70457 70556: gap of 100 bp
* 70557 71570: contig of 1014 bp in length
* 71571 71670: gap of 100 bp
* 71671 72655: contig of 985 bp in length.

Location/Qualifiers

1. 72655
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"

FEATURES
source

Query Match 84.8%; Score 17.8; DB 2; Length 72655;
Best Local Similarity 90.5%; Pred. No. 4.5e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTAGCT 21
 Db 26318 CTGTGACATGCAGATCTAGCT 26298

RESULT 8
 LOCUS AL831718/c
 DEFINITION Mouse DNA sequence from clone RP23-146020 on chromosome X, complete sequence.
 ACCESSION AL831718
 VERSION AL831718.6 GI:25137019
 KEYWORDS HTG.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 116076)
 AUTHORS Clark, S.
 TITLE Direct Submission
 JOURNAL Submitted (13-AUG-2002) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
 COMMENT On Nov 19, 2002 this sequence version replaced gi:22213735.
 ----- Genome Center
 Center: Wellcome Trust Sanger Institute
 Center code: SC
 Web site: <http://www.sanger.ac.uk>
 Contact: humquery@sanger.ac.uk

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.
 This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TrEMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep RP23-146020 is from the RPCI-23 Mouse PAC library
 constructed by the group of Pieter de Jong.
 For further details see <http://www.chori.org/bacpac/home.htm>
 VECTOR: pBAC3.6.

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 /chromosome="X"
 /clone="RP23-146020"
 /clone_lib="RPCI-23"

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 Best Local Similarity 90.5%; Pred. No. 4.2e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTAGCT 21
 Db 107376 CTGTGACATGCAGATCTAGCT 107356

RESULT 9
 LOCUS AC148761/c
 DEFINITION

Medicago truncatula chromosome 2 clone mth2-19h23, *** SEQUENCING IN PROGRESS ***, 14 unordered pieces.
 AC148761

ACCESSION AC148761.1
 VERSION GI:46063628
 KEYWORDS HTG; HTGS PHASE1; HTGS ACTIVEPIN.
 SOURCE Medicago truncatula (barrel medic)
 ORGANISM Medicago truncatula

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Trifolieae; Medicago.

REFERENCE 1 (bases 1 to 125354)
 AUTHORS Town, C.D., Tallon, L.J., Arbogast, T., Althoff, R., Hine, E., Monaghan, E., Smith, S.A., Utterback, T., Feldblyum, T. and Koo, H.
 TITLE Medicago truncatula BAC genomic sequence
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 125354)
 AUTHORS Town, C.D.

Direct Submission
 Submitted (02-APR-2004) The Institute for Genomic Research, 9712 Medical Center Dr, Rockville, MD 20850, USA

NOTE: This is a 'working draft' sequence. It currently consists of 14 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 1075: contig of 1075 bp in length
 * 1076 1175: gap of unknown length
 * 1176 2410: contig of 1235 bp in length
 * 2411 2510: gap of unknown length
 * 2511 29548: contig of 27038 bp in length
 * 29549 29648: gap of unknown length
 * 29649 32369: contig of 2721 bp in length
 * 32370 32469: gap of unknown length
 * 32470 44227: contig of 11758 bp in length
 * 44228 44327: gap of unknown length
 * 44328 45752: contig of 1425 bp in length
 * 45753 45852: gap of unknown length
 * 45853 77661: contig of 31809 bp in length
 * 77662 77761: gap of unknown length
 * 77762 79043: contig of 1282 bp in length
 * 79044 88662: contig of 9519 bp in length
 * 88663 88762: gap of unknown length
 * 88763 91773: contig of 3011 bp in length
 * 91774 91873: gap of unknown length
 * 91874 95698: contig of 3825 bp in length
 * 95699 95798: gap of unknown length
 * 95799 103010: contig of 7212 bp in length
 * 103011 103110: gap of unknown length
 * 103111 111029: contig of 7919 bp in length
 * 11030 111129: gap of unknown length
 * 111130 125354: contig of 14225 bp in length.

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Query Match 84.8%; Score 17.8; DB 2; Length 125354;
 Best Local Similarity 90.5%; Pred. No. 4.2e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGCTAGCT 21

Db 42736 CAGTGACATGAGCGCTACCT 42716

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RESULT 10
AC110379 AC110379 139884 bp DNA linear ROD 05-NOV-2003
LOCUS Mus musculus BAC clone RP24-193N24 from 15, complete sequence.
ACCESSION AC110379
VERSION AC110379.3 GI:21536156
KEYWORDS HTG.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 139884)
Haakenson, W. and Doeber, A.
The sequence of Mus musculus BAC clone RP24-193N24
Unpublished (2001)
REFERENCE 2 (bases 1 to 139884)
AUTHORS Wilson, R.
Sequencing of Mus musculus
Unpublished (2001)
REFERENCE 3 (bases 1 to 139884)
AUTHORS McPherson, J.D. and Waterston, R.H.
Direct Submission
TITLE Submitted (11-FEB-2002) Genome Sequencing Center, 4444 Forest Park
JOURNAL Parkway, St. Louis, MO 63108, USA
4 (bases 1 to 139884)
McPherson, J.D. and Waterston, R.H.
Direct Submission
TITLE Submitted (08-APR-2002) Genome Sequencing Center, 4444 Forest Park
JOURNAL Parkway, St. Louis, MO 63108, USA
5 (bases 1 to 139884)
McPherson, J.D. and Waterston, R.H.
Direct Submission
TITLE Submitted (21-JUN-2002) Genome Sequencing Center, 4444 Forest Park
JOURNAL Parkway, St. Louis, MO 63108, USA
6 (bases 1 to 139884)
Wilson, R.
Direct Submission
TITLE Submitted (05-NOV-2003) Department of Genetics, Washington
JOURNAL University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Jun 21, 2002 this sequence version replaced gi:20069820.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu
Contact: submissions@watson.wustl.edu
----- Summary Statistics
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Center project name: M_BB0193N24
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NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. Wes Warren, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see http://genome.wustl.edu

SOURCE INFORMATION:

NEIGHBORING SEQUENCE INFORMATION:

This sequence is the entire insert of the clone.

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| | /mol_type="genomic DNA" |
| | /db_xref="taxon:10090" |
| | /chromosome="15" |
| | /map="15" |
| | /clone="RP24-193N24" |
| | /clone_lib="RPCI-24" |
| repeat_region | 1472..1532 |
| | /rpt_family="Alu" |
| repeat_region | 2336..2478 |
| | /rpt_family="B4" |
| repeat_region | 3625..3939 |
| | /rpt_family="MaLR" |
| repeat_region | 4251..4445 |
| | /rpt_family="B2" |
| repeat_region | 4623..4984 |
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| repeat_region | 6912..7089 |
| | /rpt_family="B2" |
| repeat_region | 7105..7222 |
| | /rpt_family="B4" |
| repeat_region | 7264..7331 |
| | /rpt_family="Alu" |
| repeat_region | 7640..7961 |
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| | /rpt_family="B4" |
| repeat_region | 11045..11288 |
| | /rpt_family="MaLR" |
| repeat_region | 11701..11779 |
| | /rpt_family="L1" |
| repeat_region | 12001..12197 |
| | /rpt_family="B2" |
| repeat_region | 12903..13063 |
| | /rpt_family="MER1_type" |
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| | /rpt_family="B2" |
| repeat_region | 14427..14547 |
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| | /rpt_family="MaLR" |
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| | /rpt_family="L1" |
| repeat_region | 18215..18343 |
| | /rpt_family="L1" |
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| repeat_region | 21199..21262 |
| | /rpt_family="4.5SRNA" |
| repeat_region | 21210..21311 |
| | /rpt_family="Alu" |
| repeat_region | 21500..22334 |
| | /rpt_family="L1" |
| repeat_region | 22533..22620 |

The RPCI-24 BAC Library has been constructed by Pieter de Jong and coworkers (http://www.chori.org) from male C57BL/6J mouse spleen and/or brain genomic DNA. The clone and detailed information can be obtained from Pieter de Jong and coworkers at http://www.chori.org

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* 8147 9901: contig of 1755 bp in length
* 9902 10001: gap of unknown length
* 10002 12500: contig of 2499 bp in length
* 12501 12600: gap of unknown length
* 12601 15266: contig of 2666 bp in length
* 15267 15366: gap of unknown length
* 15367 17549: contig of 2183 bp in length
* 17550 20378: gap of unknown length
* 20379 20478: gap of unknown length
* 20479 23345: contig of 2867 bp in length
* 23346 27122: contig of 3677 bp in length
* 27123 30048: gap of unknown length
* 30049 30148: gap of unknown length
* 30149 32220: contig of 2072 bp in length
* 32221 32320: gap of unknown length
* 32321 35967: contig of 3647 bp in length
* 35968 36067: gap of unknown length
* 36068 39730: contig of 3663 bp in length
* 39731 39830: gap of unknown length
* 39831 43447: contig of 3617 bp in length
* 43448 43547: gap of unknown length
* 43548 47467: contig of 3920 bp in length
* 47468 47567: gap of unknown length
* 47568 51880: contig of 4313 bp in length
* 51881 51980: gap of unknown length
* 51982 57315: contig of 5335 bp in length
* 57316 57415: gap of unknown length
* 57416 61833: contig of 4418 bp in length
* 61834 61933: gap of unknown length
* 61934 65951: contig of 4018 bp in length
* 65952 66051: gap of unknown length
* 66052 70863: contig of 4818 bp in length
* 70864 70963: gap of unknown length
* 70964 77112: contig of 6043 bp in length
* 77113 77113: gap of unknown length
* 77114 82785: contig of 5673 bp in length
* 82786 82885: gap of unknown length
* 82886 88405: contig of 5520 bp in length
* 88406 88505: gap of unknown length
* 88506 93767: contig of 5262 bp in length
* 93768 93867: gap of unknown length
* 93868 99572: contig of 5705 bp in length
* 99573 99672: gap of unknown length
* 99673 109490: contig of 9818 bp in length
* 109491 109590: gap of unknown length
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* 131548 131647: gap of unknown length
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* 148259 148358: gap of unknown length
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ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 162560;
Best Local Similarity 90.5%; Pred. No. 4.1e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGCTTAGCT 21

Db 64559 CAGTGACATGCAGCTTAGCT 64539

RESULT 14
AC103719/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

AC103719 167878 bp DNA linear PRI 07-JAN-2003
Homo sapiens chromosome 8, clone RP11-421P23, complete sequence.
AC103719
AC103719.12 GI:27531859
HTG.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 167878)
Homo sapiens chromosome 8, clone RP11-421P23
Unpublished
2 (bases 1 to 167878)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,
Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Bouckgalter, B.,
Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B.,
Choepe, Y., Collangelo, M., Collins, S., Collymore, A., Cook, A.,
Cooke, P., DeArellano, K., Dewar, K., Diaz, J. S., Dodge, S., Fato, S.,
Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S.,
Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,
Hagos, B., Hearford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,
Jones, C., Kamat, A., Karatas, A., Kells, C., Larocque, K.,
Lamazares, R., Landers, T., Lehoczy, J., Levine, R., Liu, G.,
MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C.,
McCarthy, M., McEwan, P., McKernan, K., McPheeters, R., Meldrum, J.,
Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C.,
Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D.,
Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollard, V.,
Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,
Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupback, R.,
Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
Topham, K., Travers, M., Travis, N., Triggillo, J., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G.,
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission

Submitted (29-NOV-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 167878)

JOURNAL

REFERENCE

AUTHORS

Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Bouckgalter, B.,
Camarata, J., Chang, J., Chazaro, B., Choepe, Y., Collymore, A.,
Cooke, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J. S., Dodge, S.,
Fato, S., Ferreira, P., FitzGerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagel, N.,
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R.,
Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J.,
Matthews, C., McCarthy, M., Meldrum, J., Meneus, L., Mihova, T.,
Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,
Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,
Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R.,
Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schupback, R.,
Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N.,
Stojanovic, N., Talamas, J., Tesfaye, S., Theodore, J., Topham, K.,
Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X.,
Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (03-JAN-2003) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 167878)

JOURNAL

REFERENCE

AUTHORS

Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Bouckgalter, B.,
Camarata, J., Chang, J., Chazaro, B., Choepe, Y., Collymore, A.,
Cooke, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J. S., Dodge, S.,
Fato, S., Ferreira, P., FitzGerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagel, N.,
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R.,


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KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
FEATURES
source

HTG.
Homo sapiens (human)
Homo sapiens
Mammalia; Eutheria; Chordata; Craniata; Vertebrata; Euteleostomi;
Eukaryota; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 189662)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Homo sapiens chromosome 8, clone RP11-369E15
Unpublished
2 (bases 1 to 189662)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Baldwin,J., Barna,N., Beckerly,R., Boguslavskiy,L., Bouckgalter,B.,
Brown,A., Castle,A., Colangelo,M., Collins,S., Collymore,A.,
Cooke,P., Dearellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,
Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,
Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Lehoczky,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,
Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severi,P.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
Wyman,D., Ye.W.J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (16-NOV-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 189662)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Boguslavskiy,L., Bouckgalter,B., Brown,A.,
Camarata,J., Campopiano,A., Chang,J., Choepel,Y., Colangelo,M.,
Collins,S., Collymore,A., Cooke,P., Dearellano,K., Dewar,K.,
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Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C.,
Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J.,
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Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,
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Theodore,J., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye.W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (01-MAY-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On May 1, 2001 this sequence version replaced gi:12313808.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L2466
Center clone name: 369_E_15
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/clone_lib="RPC1-11 Human Male BAC"
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complement(16078. .16165)
/rpt family="AluJ/FRAM"
16532. .17094
/rpt family="MLT1F1"
17653. .17674
/rpt family="(TAA)n"
complement(17675. .17892)
/rpt family="MER7C"
complement(17897. .18650)
/rpt family="LTR17"
complement(18651. .18996)
/rpt family="MER7C"
18999. .19103
/rpt family="(TAA)n"
complement(19203. .19283)
/rpt family="MIR"

```


contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
Center: Baylor College of Medicine

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: KATP

Center clone name: CH230-304B3

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 191737 bases at least Q40

Consensus quality: 193479 bases at least Q30

Consensus quality: 194485 bases at least Q20

Estimated insert size: 194733; sum-of-contigs estimation

Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 5 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 191071: contig of 191071 bp in length

* 191072 191171: gap of unknown length

* 191172 192706: contig of 1535 bp in length

* 192707 192806: gap of unknown length

* 192807 194126: contig of 1320 bp in length

* 194127 194226: gap of unknown length

* 194227 196128: contig of 1902 bp in length

* 196129 196228: gap of unknown length

* 196229 197796: contig of 1568 bp in length.

* Location/Qualifiers

1..197796

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-304B3"

1..1063

/note="wgs_end_extension

clone_end:17"

1565..2730

/note="wgs_end_extension

clone_end:17"

3873..4716

/note="clone_boundary

clone_end:17

site:

end_sequence:BZ204887"

ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 197796;

Best Local Similarity 90.5%; Pred. No. 4e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGTGACATGACGGTCTAGCT 21

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VERSION

KEYWORDS

SOURCE

ORGANISM

AC129792.4 GI:25073629

HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.

Rattus norvegicus (Norway rat)

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 197796)

REFERENCE

AUTHORS

Muzny,D.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Blewato,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,Y., Chen,Y., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gebregiorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,M.,
Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hogues,M.,
Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpachy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensuhera,L., Loulseghe,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munitasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwaokemele,O., Okwuonu,G., Olarnpusagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkuch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.,
Puzo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.,
Sanders,W., Savery,G., Scherer,S., Scott,G., Shatman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sison,I., Sitter,C.D., Smajs,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Umami,K.,
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,
Wang,O., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wlezyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs.R.A.

Direct Submission

Unpublished

2 (bases 1 to 197796)

Worley,K.C.

Direct Submission

Submitted (03-AUG-2002)

Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 197796)

Rat Genome Sequencing Consortium.

Direct Submission

Submitted (19-NOV-2002)

Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

On Nov 19, 2002 this sequence version replaced gi:23915295.

The sequence in this assembly is a combination of BAC based reads

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: KATP
Center clone name: CH230-304B3

----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 191737 bases at least Q40
Consensus quality: 193479 bases at least Q30
Consensus quality: 194485 bases at least Q20
Estimated insert size: 194733; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently consists of 5 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

* 1 191071: contig of 191071 bp in length
* 191072 191171: gap of unknown length
* 191172 192706: contig of 1535 bp in length
* 192707 192806: gap of unknown length
* 192807 194126: contig of 1320 bp in length
* 194127 194226: gap of unknown length
* 194227 196128: contig of 1902 bp in length
* 196129 196228: gap of unknown length
* 196229 197796: contig of 1568 bp in length.

FEATURES
source
1..197796
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-304B3"
1..1063
/note="wgs_end_extension
clone_end:77"
1565..2730
/note="wgs_end_extension
clone_end:77"
3873..4716
/note="clone_boundary
clone_end:77
site:
end_sequence:BZ204887"

ORIGIN
Query Match 84.8%; Score 17.8; DB 2; Length 197796;
Best Local Similarity 90.5%; Pred. No. 4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 CAGTCACATGCGGCTAGCT 21
DB 97483 CTGTGATATGACGGTCTAGCT 97463

RESULT 18

AC087221

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

AC087221 203690 bp DNA linear HTG 24-MAY-2002
Homo sapiens chromosome 8 clone RP11-712115 map 8, WORKING DRAFT
SEQUENCE, 34 ordered pieces.
AC087221
HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_FULLTOP.
Homo sapiens (human)

AC087221.2 GI:21166223
HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_FULLTOP.
Homo sapiens
Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 203690)
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 8, clone RP11-712115
Unpublished
2 (bases 1 to 203690)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B., Brown,A.,
Camarata,J., Campopiano,A., Choepel,Y., Colangelo,M., Collins,S.,
Collins,S., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Faro,S., Ferreira,P., FitzHugh,W., Gage,D., Galagan,J.,
Gardyna,S., Ginde,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Karatas,A., LaRocque,K., Lamazares,R., Landers,T.,
Lehoczy,J., Levine,R., Liu,G., Maclean,C., Macdonald,P.,
Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K.,
McPheeters,R., Meldrim,J., Meneus,L., Mihova,T., Miengo,V.,
Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R.,
Rieback,M., Riley,P., Rise,C., Rogov,P., Roman,J., Rosetti,M.,
Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S., Severy,P.,
Sougnaz,C., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Testfaye,S., Theodore,J.,
Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo,A.,
Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.

Direct Submission
Submitted (16-DEC-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 203690)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L.,
Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,
Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A.,
Cooke,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I.,
Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,
Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Marquis,N.,
Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrim,J.,
Meneus,L., Mihova,T., Miengo,V., Murphy,T., Naylor,J., Nguyen,C.,
Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C.,
Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S.,
Schupback,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testfaye,S.,
Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission
Submitted (24-MAY-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On May 24, 2002 this sequence version replaced gi:11875303.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L11638

Center clone name: 712_1_15

----- Summary Statistics

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 190084 bases at least Q40

Consensus quality: 196705 bases at least Q30

Consensus quality: 199369 bases at least Q20

Insert size: 176000; agarose-fp

Insert size: 200390; sum-of-contigs

Quality coverage: 9.6 in Q20 bases; agarose-fp

Quality coverage: 8.5 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 34 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* been provided by the submitter.

* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

* 1 460: contig of 460 bp in length

* 461 560: gap of 100 bp

* 561 1196: contig of 636 bp in length

* 1197 1296: gap of 100 bp

* 1297 2122: contig of 826 bp in length

* 2123 2222: gap of 100 bp

* 2223 2869: contig of 647 bp in length

* 2870 2969: gap of 100 bp

* 2970 3620: contig of 651 bp in length

* 3621 3720: gap of 100 bp

* 3721 4314: contig of 594 bp in length

* 4315 4414: gap of 100 bp

* 4415 5152: contig of 738 bp in length

* 5153 5252: gap of 100 bp

* 5253 6209: contig of 957 bp in length

* 6210 6309: gap of 100 bp

* 6310 7103: contig of 794 bp in length

* 7104 7203: gap of 100 bp

* 7204 7937: contig of 734 bp in length

* 7938 8037: gap of 100 bp

* 8038 9006: contig of 969 bp in length

* 9007 9106: gap of 100 bp

* 9107 9906: contig of 800 bp in length

* 9907 10006: gap of 100 bp

* 10007 11070: contig of 1064 bp in length

* 11071 11170: gap of 100 bp

* 11171 12297: contig of 1127 bp in length

* 12298 12397: gap of 100 bp

* 12398 13415: contig of 1018 bp in length

* 13416 13515: gap of 100 bp

* 13516 14231: contig of 716 bp in length

* 14232 14331: gap of 100 bp

* 14332 15596: contig of 1265 bp in length

* 15597 15696: gap of 100 bp

* 15697 16462: contig of 766 bp in length

* 16463 16562: gap of 100 bp

* 16563 17742: contig of 1180 bp in length

* 17743 17842: gap of 100 bp

* 17843 18857: contig of 1015 bp in length

* 18858 18957: gap of 100 bp

* 18958 20490: contig of 1533 bp in length

* 20491 20590: gap of 100 bp

* 20591 22210: contig of 1820 bp in length

* 22211 22310: gap of 100 bp

* 22311 23851: contig of 1541 bp in length

* 23852 23951: gap of 100 bp

* 23952 25684: contig of 1733 bp in length

* 25685 25784: gap of 100 bp

* 25785 27006: contig of 1222 bp in length

* 27007 27106: gap of 100 bp

* 27107 28616: contig of 1510 bp in length

* 28617 28716: gap of 100 bp

* 28717 30184: contig of 1468 bp in length

* 30185 30284: gap of 100 bp

* 30285 31303: contig of 1019 bp in length

* 31304 31403: gap of 100 bp

* 31404 32917: contig of 1514 bp in length

* 32918 33017: gap of 100 bp

* 33018 42030: contig of 9013 bp in length

* 42031 42130: gap of 100 bp

* 42131 49012: contig of 6882 bp in length

* 49013 49112: gap of 100 bp

* 49113 62240: contig of 13128 bp in length

* 62241 62340: gap of 100 bp

* 62341 86795: contig of 24455 bp in length

* 86796 86895: gap of 100 bp

* 86896 203690: contig of 116795 bp in length.

FEATURES

source

Location/Qualifiers

1..203690

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/chromosome="8"

/map="8"

/clone="RP11-712115"

/clone_lib="RPC1-11 Human Male BAC"

1..460

misc_feature

/note="assembly_fragment"

clone end:SP6

vector side:left"

561..1196

/note="assembly_fragment"

1297..2122

/note="assembly_fragment"

2223..2869

/note="assembly_fragment"

2970..3620

/note="assembly_fragment"

3721..4314

/note="assembly_fragment"

4415..5152

/note="assembly_fragment"

5253..6209

/note="assembly_fragment"

6310..7103

/note="assembly_fragment"

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/note="assembly_fragment"

8038..9006

/note="assembly_fragment"

9107..9906

/note="assembly_fragment"

Query Match 84.8%; Score 17.8; DB 2; Length 203690;

Best Local Similarity 90.5%; Pred.No.4e+02; 2; Indels 0; Gaps 0;

Matches 19; Conservative 0; Mismatches 2;

Qy 1 CAGTGACATGCAGGCTCTAGCT 21

|||||

Db 190273 CAGTGACATGCAGGCTCTAGCT 190293

RESULT 19

AC115746

LOCUS

AC115746

DEFINITION Mus musculus chromosome 15, clone RP23-3J8, complete sequence.

ACCESSION AC115746

VERSION AC115746.10 GI:50811761

214765 bp DNA linear ROD 29-JUL-2004


```

repeat_region 12370. 12444
/rpt family="CA)n"
repeat_region 13343. 13367
/rpt family="TGGGG)n"
repeat_region 14362. 14384
/rpt family="AT rich"
repeat_region complement(15620. 15776)
/rpt family="B1_MM"
repeat_region 15792. 15822
/rpt family="TTTC)n"
repeat_region complement(15830. 15977)
/rpt family="B1_MM"
repeat_region complement(16486. 16564)
/rpt family="ID1_MM"
repeat_region complement(17181. 17277)
/rpt family="RSINE1"
repeat_region 17293. 17388
/rpt family="B4"
repeat_region 17391. 17584
/rpt family="B2_Mm2"
repeat_region 17585. 17621
/rpt family="polypurine"
repeat_region 17623. 17960
/rpt family="(GGA)n"
repeat_region 18167. 18196
/rpt family="(CA)n"
repeat_region 18533. 18595
/rpt family="ID5"
repeat_region 18596. 18608
/rpt family="ID_B1"
repeat_region 19265. 19413
/rpt family="B1_MM"
repeat_region 19715. 19921
/rpt family="B4A"
repeat_region 19952. 19992
/rpt family="(CA)n"
repeat_region 20016. 20141
/rpt family="PB1D9"
repeat_region 20286. 20398
/rpt family="B1_MM"
repeat_region 20399. 20488
/rpt family="GA-rich"
repeat_region 20736. 20898
/rpt family="B3A"
repeat_region 20931. 20957
/rpt family="(CA)n"
repeat_region 22176. 22256
/rpt family="(GA)n"
repeat_region 23275. 23413
/rpt family="B1_MM"

Query Match 84.8%; Score 17.8; DB 10; Length 214765;
Best Local Similarity 90.5%; Pred.No. 3.9e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTAGCT 21
Db 98096 CAGTGACTTGCAGGCTAGCT 98116
|||||
|||||

RESULT 20
AC120123/c
LOCUS AC120123
DEFINITION Mus musculus chromosome 7 clone RP23-152B12 map 7, *** SEQUENCING
IN PROGRESS ***, 7 unordered pieces.
ACCESSION AC120123
VERSION AC120123.10 GI:52139883
KEYWORDS HTG; HTGS PHASE1; HTGS FULLTOP; HTGS_ACTIVEPIN.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 222540)
Birren,B., Nusbaum,C. and Lander,E.
Mus musculus chromosome 7, clone RP23-152B12
Unpublished
2 (bases 1 to 222540)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L.,
Bouhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,
Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N., Jones,C.,
Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kamat,A., Karatas,A., Kells,C., LaRocque,K., Lamazares,R.,
Landers,T., Lehoczyk,J., Levine,R., Lindblad-Toh,K., Liu,G.,
MacLean,C., MacDonald,P., Major,J., Marguis,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., Meldrim,J., Meneus,L.,
Mihova,T., Mlenka,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (03-MAY-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 222540)
Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,
Anderson,M., Anderson,S., Arachchi,H.M., Barna,N., Bastien,V.,
Bloom,T., Boguslavskiy,L., Bouhgalter,B., Camarata,J., Chang,J.,
Choepel,Y., Collymore,A., Cook,A., Cooke,P., Corum,B.,
Dearellano,K., Diaz,J.S., Dodge,S., Dooley,K., Dorris,L.,
Erickson,J., Faro,S., Ferreira,P., FitzGerald,M., Gage,D.,
Galagan,J., Gardyna,S., Graham,L., Grand-Pierre,N., Hafez,N.,
Hagopian,D., Hagos,B., Hall,J., Horton,L., Hulme,W., Iliev,I.,
Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T.,
Levine,R., Lindblad-Toh,K., Liu,G., Liu,X., Lui,A., Mabbitt,R.,
MacLean,C., MacDonald,P., Major,J., Manning,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenka,V.,
Murphy,T., Naylor,J., Nguyen,C., Nguyen,T., Nicol,R., Norbu.C.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Rachupka,A., Ramasamy,U., Raymond,C.,
Retta,R., Rise,C., Rogov,P., Roman,J., Schauer,S., Schupback,R.,
Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Stubbs,M., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Vassiliev,H., Venkataraman,V.S., Viel,R.,
Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (16-SEP-2004) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 16, 2004 this sequence version replaced gi:50284650.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/MIT Center for Genome Research
Center code: WIRK
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@broad.mit.edu
----- Project Information
Center project name: L15655
Center clone name: 152_B_12
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 7 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence

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* as soon as it is available and the accession number will
* be preserved.
*
* 47731: contig of 47731 bp in length
* 47732: gap of unknown length
* 47832: contig of 13891 bp in length
* 61722: gap of unknown length
* 61823: contig of 91308 bp in length
* 153130: gap of unknown length
* 153231: contig of 17322 bp in length
* 170552: gap of unknown length
* 170553: contig of 20175 bp in length
* 190828: gap of unknown length
* 190928: contig of 25691 bp in length
* 216619: gap of unknown length
* 216719: contig of 5822 bp in length.
FEATURES
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    /map="7"
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    /clone_lib="RPCI-23 Female Mouse BAC"
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    Best Local Similarity 90.5%; Pred. No. 3.9e+02;
    Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 CAGTGACATGCAGGCTAGCT 21
    ||||| ||||| ||||| |||||
Db 177745 CAGTGATGACAGGCTAGCT 177725
RESULT 21
AC115307
LOCUS AC115307 260600 bp DNA linear HTG 09-NOV-2002
DEFINITION Rattus norvegicus clone CH230-11F18, WORKING DRAFT SEQUENCE.
ACCESSION AC115307
VERSION AC115307.4 GI:24817861
KEYWORDS HTG; HTGS, PHASE2; HTGS, DRAFT; HTGS, FULLTOP.
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 260600)
Muzny,D,Marie., Metzker,M, Lee., Abramzon,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Royagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,N., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Devila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gunaratne,P., Haaland,W., Hamil,C., Henderson,N., Hernandez,J.,
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hamilton,K.,
Gebregorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
Hollings,B., Howells,S., Hladun,S.L., Hodgson,A., Hogues,M.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpathy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensuewa,L., Loulseged,H., Lozado,R.J., Lu,X., Ma,J.,

```

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Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhiney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwackelmech,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfankoch,C.,
Plopper,P., Poindexter,A., Popovic,D., Primus,E., Pu,L.-L.,
Puzo,M., Quiroz,J., Rachin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Umani,K.,
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wleczyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 260600)
Worley,K.C.
Direct Submission
Submitted (17-MAR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 260600)
Rat Genome Sequencing Consortium.
Direct Submission
Unpublished
On Nov 9, 2002 this sequence version replaced gi:22772493.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GEBE
Center clone name: CH230-11F18
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 229014 bases at least Q40
Consensus quality: 231787 bases at least Q30
Consensus quality: 232904 bases at least Q20
Estimated insert size: 237320; sum-of-contigs estimation
Quality coverage: 7x in Q20 bases; sum-of-contigs estimation
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* NOTE: Estimated insert size may differ from sequence length
(fsee http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
consists of 1 contigs. Gaps between the contigs
are represented as runs of N. The order of the pieces
is believed to be correct as given, however the sizes

```

* of the gaps between them are based on estimates that have
 * provided by the submittor.
 * This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.
 * 1 260600: contig of 260600 bp in length.

FEATURES

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site:EcoRI
end_sequence:BH340449"
258937..260600
/note="wgs_end_extension"
clone_end:Sp6"

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Best Local Similarity 90.5%; Pred. NO. 3.9e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGACGAGTCTAGCT 21
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Db 178018 CAGTGACATGAGGTCTTGT 178038

RESULT 22
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LOCUS
DEFINITION Rattus norvegicus clone CH230-233D15, *** SEQUENCING IN PROGRESS ***
AC129380
VERSION AC129380.3 GI:30578573
KEYWORDS HTG: HTGS_PHASE2; HTGS_DRAFT; HTGS_ENRICHED.
SOURCE Rattus norvegicus
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 281447)
Muzny,D.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
Allen,C., Allien,H., Albrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswalo,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Detamo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Georgopoulos,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.B., Hodgson,A., Hogues,M.,
Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,

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Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpathy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kovis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensuhewa,L., Loulseghe,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhinney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minje,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Muidasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwaokaleh,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfankuch,C.,
Plopper,F., Poindestre,A., Popovic,D., Primus,E., Pu.L.-L.,
Puazo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savary,G., Scherer,S., Scott,G., Shatman,S., Shen,H.,
Shetty,J., Shvartsbeyn,A., Sison,I., Sitter,C.D., Smajd,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steimle,M., Strong,R., Sutton,A., Svatek,A., Taber,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wlezyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 281447)
Worley,K.C.
Direct Submission
Submitted (29-JUL-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 281447)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (13-MAY-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On May 13, 2003 this sequence version replaced gi:23116981.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GJ2M
Center clone name: CH230-233D15
----- Summary Statistics
Assembly program: Atlas 3.0;
Consensus quality: 224946 bases at least Q40
Consensus quality: 227863 bases at least Q30
Consensus quality: 229407 bases at least Q20
Estimated insert size: 234544; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation
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* NOTE: Estimated insert size may differ from sequence length

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* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 1 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces

* is believed to be correct as given, however the sizes

* of the gaps between them are based on estimates that have

* provided by the submittor.

* This sequence will be replaced

* by the finished sequence as soon as it is available and

* the accession number will be preserved.

* 1 281447: contig of 281447 bp in length.

FEATURES

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/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

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/note="clone boundary

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site:ECORI

end sequence:B2091967"

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/note="wgs end extension

clone_end:Sp6"

ORIGIN

Query Match 84.8%; Score 17.8; DB 2; Length 281447;

Best Local Similarity 90.5%; Pred. No. 3.8e+02; Indels 0; Gaps 0;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTAGCT 21

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Db 264359 CTGTGATATGCAGGCTAGCT 264339

RESULT 23

BD185177/c

LOCUS

BD185177 Novel genes and proteins encoded by the genes. PAT 17-JUN-2003

BD185177 5446 bp DNA linear

BD185177.1 GI:31877377

KEYWORDS JP 2002345493-A/20.

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 5446)

Ohara, O., Nagase, T. and Nakajima, D.

Novel genes and proteins encoded by the genes

Patent: JP 2002345493-A 20 03-DEC-2002;

KAZUSA DNA RESEARCH INSTITUTE

OS Homo sapiens (human)

PN JP 2002345493-A/20

PD 03-DEC-2002

PF 26-FEB-2002 JP 2002049046

PI OSAMU OHARA, TAKAHIRO NAGASE, DAISUKE NAKAJIMA

PC C12N15/09, C07K14/47, C07K14/54, C12N15/00

CC Novel genes and proteins encoded by the genes FH Key

Location/Qualifiers

FT CDS (2564)..(4138).

Location/Qualifiers

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/mol_type="genomic DNA"

/db_xref="taxon:9606"

ORIGIN

Query Match 82.9%; Score 17.4; DB 6; Length 5446;

Best Local Similarity 94.7%; Pred. No. 9.5e+02;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTCTAG 19

|||||

Db 833 CAGTGACAGGCGAGGCTCTAG 815

RESULT 24

AC121561/c

LOCUS

AC121561 Homo sapiens chromosome 17 clone CTD-2527B13 map 17, LOW-PASS

DEFINITION

SEQUENCE SAMPLING.

AC121561.1 GI:20986629

HTG; HTGS_PHASE0.

KEYWORDS

SOURCE

Homo sapiens

ORGANISM

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 46070)

Birren, B., Linton, L., Nusbaum, C. and Lander, E.

Homo sapiens chromosome 17, clone CTD-2527B13

Unpublished

2 (bases 1 to 46070)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,

Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L.,

Bouhagalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J.,

Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A.,

Cook, A., Cooke, P., DeArrellano, K., Dewar, K., Diaz, J.S., Dodge, S.,

Faro, S., Ferreira, P., Fitzgerald, M., FitzHugh, W., Gage, D.,

Galagan, J., Gardyna, S., Ginde, S., Gord, S., Govette, M., Graham, L.,

Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I.,

Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., LaRoque, K.,

Lamarez, R., Landers, T., Lehoczy, J., Levine, R., Lindblad-Toh, K.,

Liu, G., MacLean, C., Macdonald, P., Major, J., Marquis, N.,

Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meidrim, J.,

Meneus, L., Mihova, T., Menga, V., Murphy, T., Navlor, J., Nguyen, C.,

Nicol, R., Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P.,

O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N.,

Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C.,

Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S.,

Schuback, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N.,

Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S.,

Theodore, J., Toham, K., Travers, M., Travis, N., Trigilio, J., Ye, W.J.,

Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J.,

Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission

Submitted (20-MAY-2002) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

All repeats were identified using RepeatMasker:

Smt, A.P.A. & Green, P. (1996-1997)

<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WtBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L26773

Center clone name: 2527_E_13

* NOTE: This record contains 58 individual

* sequencing reads that have not been assembled into

* contigs. Runs of N are used to separate the reads

* and the order in which they appear is completely

* arbitrary. Low-pass sequence sampling is useful for

* identifying clones that may be gene-rich and allows

* overlap relationships among clones to be deduced.

* However, it should not be assumed that this clone

* will be sequenced to completion. In the event that

* the record is updated, the accession number will

* be preserved.

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* 1
* 684: contig of 684 bp in length
* 784: gap of 100 bp
* 1401: contig of 617 bp in length
* 1501: gap of 100 bp
* 2203: contig of 702 bp in length
* 2303: gap of 100 bp
* 2304: 2984: contig of 681 bp in length
* 3085: 3084: gap of 100 bp
* 3755: 3754: contig of 670 bp in length
* 3855: 3854: gap of 100 bp
* 4545: 4545: contig of 691 bp in length
* 4546: 4645: gap of 100 bp
* 4646: 5346: contig of 701 bp in length
* 5347: 5446: gap of 100 bp
* 5447: 6142: contig of 696 bp in length
* 6143: 6242: gap of 100 bp
* 6243: 6949: contig of 707 bp in length
* 6950: 7049: gap of 100 bp
* 7050: 7730: contig of 681 bp in length
* 7731: 7830: gap of 100 bp
* 7831: 8515: contig of 685 bp in length
* 8516: 8615: gap of 100 bp
* 8616: 9267: contig of 652 bp in length
* 9268: 9367: gap of 100 bp
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* 12473: 12572: gap of 100 bp
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* 13378: 14069: contig of 692 bp in length
* 14070: 14169: gap of 100 bp
* 14170: 14870: contig of 701 bp in length
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* 22079: 22178: gap of 100 bp
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* 26083: 26182: gap of 100 bp
* 26183: 26886: contig of 704 bp in length
* 26887: 26986: gap of 100 bp
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* 27692: 27791: gap of 100 bp
* 27792: 28494: contig of 703 bp in length
* 28594: 28594: gap of 100 bp
* 28595: 29303: contig of 709 bp in length
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* 29304 29403: gap of 100 bp
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* 30945 31640: contig of 696 bp in length
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* 38092 38191: gap of 100 bp
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* 38898 38997: gap of 100 bp
* 38998 39721: contig of 723 bp in length
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* 39821 40526: contig of 706 bp in length
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* 41403 42077: contig of 675 bp in length
* 42078 42177: gap of 100 bp
* 42178 42841: contig of 664 bp in length
* 42842 42941: gap of 100 bp
* 42942 43638: contig of 697 bp in length
* 43639 43738: gap of 100 bp
* 43739 44456: contig of 718 bp in length
* 44457 44556: gap of 100 bp
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* 45256 45355: gap of 100 bp
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FEATURES

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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="17"
/map="17"
/clone_lib="CITD2 Human BAC"
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ORIGIN

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Query Match      82.9%; Score 17.4; DB 2; Length 46070;
Best Local Similarity 94.7%; Pred. No. 7.4e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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Q/ 1 CAGTGACATGCAGGCTTAG 19

Db 31187 CAGTGACAGGCTTAG 31169

RESULT 25

```
AL133227
LOCUS      AL133227      85566 bp      DNA      linear      PRI 04-APR-2001
DEFINITION Human DNA sequence from clone RP11-394O2 on chromosome 20. Contains
            the gene for CGI-15 protein, a gene for a novel protein similar to
            KIAA0281 and Drosophila CG5336, ESTs, STSs, GSSs and a CpG island,
            complete sequence.
ACCESSION  AL133227
VERSION    AL133227.15 GI:9187135
KEYWORDS   HTG; CpG island; KIAA0281.
SOURCE     Homo sapiens (human)
```

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 85566)

REFERENCE

AUTHORS
TITLE
JOURNAL

Barlow K.
Direct Submission
Submitted (02-APR-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk

COMMENT

On Jul 14, 2000 this sequence version replaced gi:8977995.
During sequence assembly data is compared for overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
Chromosome 20, constructed by the Sanger Centre Chromosome 20
Mapping Group. Further information can be found at
<http://www.sanger.ac.uk/HGP/Chr20>
IMPORTANT: This sequence is not the entire insert of clone
RP11-39402. It may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true left end of clone RP11-39402 is at 1 in this sequence. The
true left end of clone RPS-981123 is at 85467 in this sequence. The
true right end of clone RPS-984123 is at 78450 in this sequence.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. RP11-39402 is from
the library RPCI-11.2 constructed by the group of Pieter de Jong.
For further details see
<http://www.chori.org/bacpac/home.htm>
VECTOR: pBACe3.6.

FEATURES

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Location/Qualifiers
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/chromosome="20"
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/clone_lib="RPCI-11.2"

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33..90
/note="MIR repeat: matches 107..161 of consensus"
139..438
/note="AluX repeat: matches 1..293 of consensus"
1404..1535
/note="MIR repeat: matches 54..182 of consensus"
3358..3536
/note="L2 repeat: matches 2318..2500 of consensus"

misc_feature

3438..3969
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misc_feature

3456..3976
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repeat_region

3620..3905
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3653..4183
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repeat_region

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repeat_region

6122..6156
/note="MIR repeat: matches 110..144 of consensus"

repeat_region

6577..6685
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repeat_region

7049..7178

repeat_region

/note="L2 repeat: matches 2562..2695 of consensus"
7275..7472

repeat_region

/note="MIR repeat: matches 44..254 of consensus"
7545..7917

misc_feature

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7733..8207

repeat_region

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repeat_region

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repeat_region

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8945..9114

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9536..9599

repeat_region

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misc_feature

/note="MIR repeat: matches 80..262 of consensus"
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repeat_region

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misc_feature

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misc_feature

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match: STS: Em:G52843"
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repeat_region

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repeat_region

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misc_feature

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misc_feature

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complement(14441..14659)

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16607..16778

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17746..18049

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18067..18413

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18593..18825

repeat_region

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18982..19111

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19112..19406

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19407..19518

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20312..20763

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21990..22289

repeat_region

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22377..22462

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22463..22749

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24255..24319

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26684. .26740
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26722. .26765
repeat_region /note="11 copies 4 mer catc 93% conserved"
26766. .26809
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26813. .26844
repeat_region /note="8 copies 4 mer atcc 93% conserved"
26839. .26918
repeat_region /note="4 copies 20 mer 77% conserved"
27222. .27559
repeat_region /note="LTR16A repeat: matches 95. .440 of consensus"
27583. .27748
repeat_region /note="FRAM repeat: matches 1. .166 of consensus"
27786. .27897
repeat_region /note="L2 repeat: matches 2637. .2750 of consensus"
28308. .28431
repeat_region /note="AluSg repeat: matches 1. .132 of consensus"
28448. .28524
repeat_region /note="LTR16A repeat: matches 193. .270 of consensus"
28525. .28714
repeat_region /note="AluSg repeat: matches 114. .301 of consensus"
28837. .29017
repeat_region /note="MIR repeat: matches 65. .250 of consensus"
29053. .29187
repeat_region /note="MIR repeat: matches 84. .226 of consensus"
30560. .31063
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30596. .30931
misc_feature /note="match: GSS: Em:AQ901120"
30602. .30764
repeat_region /note="MER58A repeat: matches 46. .221 of consensus"
complement(31347. .46204)
gene /gene="BA39402.1"
mRNA complement(join(31347. .32333,32570. .32699,33860. .34035,
36691. .36774,36882. .37050,37611. .37683,38383. .38446,
39432. .39582,40196. .40572,46180. .46204))
/gene="BA39402.1"
/product="BA39402.1 (CGI-15 protein)"
/note="match: cDNAs: Em:AF132949
match: ESTs: Em:AU079917 Em:AA667893 Em:AI007286
Em:AU067617 Em:W56183 Em:R69763 Em:H06603 Em:AW631237"

Query Match 82.9%; Score 17.4; DB 9; Length 85566;
Best Local Similarity 94.7%; Pred. No. 6.9e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CAGTCACATGCAGGTCTAG 19
Db 64014 CAGTCACAGGCAGGTCTAG 64032
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Search completed: September 6, 2005, 20:29:56
Job time : 746.656 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 189.656 Seconds
(without alignments)
655.473 Million cell updates/sec

Title: US-10-729-421-40
Perfect score: 21
Sequence: 1 cagtgcacatgcagcttagct 21

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 100 summaries

Database : N_Geneseq_16Dec04.*
1: geneseqn1980s.*
2: geneseqn1990s.*
3: geneseqn2000s.*
4: geneseqn2001as.*
5: geneseqn2001bs.*
6: geneseqn2002as.*
7: geneseqn2002bs.*
8: geneseqn2003as.*
9: geneseqn2003bs.*
10: geneseqn2003cs.*
11: geneseqn2003ds.*
12: geneseqn2004as.*
13: geneseqn2004bs.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

| Result | No. | Score | Query | Match | Length | ID | Description |
|--------|-----|-------|-------|-------|--------|----------|--------------------|
| C | 1 | 21 | 100.0 | 21 | 8 | ACC80536 | Acc80536 Exemplary |
| C | 2 | 21 | 100.0 | 21 | 8 | ACC80538 | Acc80538 Internal |
| C | 3 | 21 | 100.0 | 21 | 8 | ACC80537 | Acc80537 Internal |
| C | 4 | 21 | 100.0 | 21 | 9 | ABZ59632 | ABZ59632 Parvoviru |
| C | 5 | 21 | 100.0 | 21 | 12 | ADI53796 | ADI53796 HAV inter |
| C | 6 | 21 | 100.0 | 23 | 12 | ADI53797 | ADI53797 HAV inter |
| C | 7 | 21 | 100.0 | 23 | 12 | ADQ30670 | Adq30670 West Nile |
| C | 8 | 21 | 100.0 | 25 | 12 | ADQ30671 | Adq30671 West Nile |
| C | 9 | 21 | 100.0 | 681 | 9 | ABZ59634 | ABZ59634 Exemplary |
| C | 10 | 21 | 100.0 | 727 | 12 | ADI53795 | ADI53795 HAV inter |
| C | 11 | 21 | 100.0 | 967 | 12 | ADQ30647 | Adq30647 West Nile |
| C | 12 | 21 | 100.0 | 1696 | 8 | ACC80539 | Acc80539 Internal |
| C | 13 | 17.4 | 82.9 | 5446 | 10 | ADE71204 | Ade71204 Novel hum |
| C | 14 | 17.4 | 82.9 | 28482 | 8 | ABZ73855 | Abz73855 Secreted |
| C | 15 | 17.4 | 82.9 | 28482 | 8 | ADA44262 | Ada44262 Human sec |
| C | 16 | 17.4 | 82.9 | 32681 | 8 | ABZ74517 | Abz74517 Secreted |
| C | 17 | 17.4 | 82.9 | 32681 | 8 | ABZ73854 | Abz73854 Secreted |
| C | 18 | 17.4 | 82.9 | 32681 | 8 | ADA98915 | Ada98915 Human sec |
| C | 19 | 17.4 | 82.9 | 32681 | 8 | ADA44261 | Ada44261 Human sec |
| C | 20 | 17.4 | 82.9 | 32681 | 8 | ADA44519 | Ada44519 Human sec |

| | | | | | | | |
|---|----|------|------|--------|----|------------|--------------------|
| C | 21 | 17.4 | 82.9 | 32681 | 10 | ADC20949 | Adc20949 Human sec |
| C | 22 | 17.4 | 82.9 | 32681 | 10 | ABZ68053 | Abz68053 Human sec |
| C | 23 | 16.8 | 80.0 | 320 | 8 | ABX46275 | Abx46275 Bovine ES |
| C | 24 | 16.8 | 80.0 | 36221 | 4 | AAS00624 | Aas00624 Human Gaa |
| C | 25 | 16.4 | 78.1 | 72705 | 11 | ACN45158 | Acn45158 Human Gen |
| C | 26 | 16.4 | 78.1 | 110000 | 8 | ABX16390_5 | Continuation (6 of |
| C | 27 | 16.4 | 78.1 | 117382 | 11 | ACN44804 | Acn44804 Mouse gen |
| C | 28 | 16.4 | 78.1 | 340449 | 8 | AAL52198 | Aal52198 Human sec |
| C | 29 | 16.2 | 77.1 | 201 | 13 | ADS40801 | Ads40801 Human aut |
| C | 30 | 16.2 | 77.1 | 201 | 13 | ADS39530 | Ads39530 Human aut |
| C | 31 | 16.2 | 77.1 | 394 | 4 | AAL82251 | Aal82251 Human pol |
| C | 32 | 16.2 | 77.1 | 473 | 9 | ACH36560 | Ach36560 Human end |
| C | 33 | 16.2 | 77.1 | 497 | 10 | ADB56505 | Adb56505 Toxicity- |
| C | 34 | 16.2 | 77.1 | 1487 | 10 | ADC71327 | Adc71327 Human col |
| C | 35 | 16.2 | 77.1 | 2048 | 2 | AAQ85985 | Aaq85985 zea maye |
| C | 36 | 16.2 | 77.1 | 2174 | 10 | ADBE62245 | Ade62245 Rat gene |
| C | 37 | 16.2 | 77.1 | 2238 | 5 | AAS76384 | Aas76384 DNA encod |
| C | 38 | 16.2 | 77.1 | 2569 | 13 | ADRO6803 | Adro6803 Full leng |
| C | 39 | 16.2 | 77.1 | 2721 | 8 | AAL53547 | Aal53547 cDNA of h |
| C | 40 | 16.2 | 77.1 | 4445 | 6 | ABA01096 | Aba01096 Brevibact |
| C | 41 | 16.2 | 77.1 | 9048 | 4 | AAC90812 | Aac90812 B. lactof |
| C | 42 | 16.2 | 77.1 | 10500 | 4 | AAL05334 | Aal05334 Human rep |
| C | 43 | 16.2 | 77.1 | 10500 | 4 | ABL98203 | AbL98203 Human tes |
| C | 44 | 16.2 | 77.1 | 14902 | 13 | ADS36489 | Ads36489 Human aut |
| C | 45 | 16.2 | 77.1 | 15515 | 8 | AAL53548 | Aal53548 Genomic D |
| C | 46 | 16.2 | 77.1 | 55827 | 8 | ACA60949 | AcA60949 DNA encod |
| C | 47 | 16.2 | 77.1 | 58337 | 13 | ADS36454 | Ads36454 Human aut |
| C | 48 | 16.2 | 77.1 | 64423 | 13 | ADS36462 | Ads36462 Human aut |
| C | 49 | 16.2 | 77.1 | 70372 | 6 | AAL53465 | Aal53466 Rag-like |
| C | 50 | 16.2 | 77.1 | 90442 | 9 | ADA03077 | Ada03077 Mouse mCG |
| C | 51 | 16.2 | 77.1 | 90442 | 9 | ADA66361 | Ada66361 Mouse mCG |
| C | 52 | 16.2 | 77.1 | 90442 | 10 | ADB72815 | Adb72815 Mouse mCG |
| C | 53 | 16.2 | 77.1 | 90442 | 10 | ADC26997 | Adc26997 Mouse car |
| C | 54 | 16.2 | 77.1 | 90442 | 11 | ADL27155 | AdL27155 Mouse gen |
| C | 55 | 16.2 | 77.1 | 143306 | 6 | ABK49586 | Abk49586 Human tra |
| C | 56 | 16.2 | 77.1 | 349980 | 5 | AAH68529 | Aah68529 C glutami |
| C | 57 | 15.8 | 75.2 | 171 | 2 | AAV89101 | Aav89101 EST clone |
| C | 58 | 15.8 | 75.2 | 279 | 9 | AAV87960 | Aav87960 EST clone |
| C | 59 | 15.8 | 75.2 | 297 | 9 | ADB08791 | AdB08791 Alloiococ |
| C | 60 | 15.8 | 75.2 | 297 | 9 | ADB08789 | AdB08789 Alloiococ |
| C | 61 | 15.8 | 75.2 | 497 | 5 | AAS88145 | Aas88145 DNA encod |
| C | 62 | 15.8 | 75.2 | 497 | 5 | AAS80088 | Aas80088 DNA encod |
| C | 63 | 15.8 | 75.2 | 587 | 4 | AAH07183 | Aah07183 Human CDN |
| C | 64 | 15.8 | 75.2 | 748 | 4 | AAH03978 | Aah03978 Human CDN |
| C | 65 | 15.8 | 75.2 | 882 | 10 | ADC08897 | AdC08897 Rice DNA |
| C | 66 | 15.8 | 75.2 | 1614 | 9 | ADB08797 | AdB08797 Alloiococ |
| C | 67 | 15.8 | 75.2 | 1670 | 4 | AAH17173 | Aah17173 Human CDN |
| C | 68 | 15.8 | 75.2 | 1777 | 4 | AAH16391 | Aah16391 Human CDN |
| C | 69 | 15.8 | 75.2 | 1995 | 6 | ABL88382 | AbL88382 Pain regu |
| C | 70 | 15.8 | 75.2 | 2378 | 5 | AAS64699 | Aas64699 DNA encod |
| C | 71 | 15.8 | 75.2 | 2379 | 5 | AAS67041 | Aas67041 DNA encod |
| C | 72 | 15.8 | 75.2 | 4015 | 12 | ADJ34728 | Adj34728 Rat 2'-5' |
| C | 73 | 15.8 | 75.2 | 4708 | 12 | ADJ34707 | Adj34707 Mouse 2'- |
| C | 74 | 15.8 | 75.2 | 31236 | 9 | ADA02900 | Ada02900 Human PTP |
| C | 75 | 15.8 | 75.2 | 31236 | 10 | ADB72638 | Adb72638 Human PTP |
| C | 76 | 15.8 | 75.2 | 31236 | 10 | ADC85379 | Adc85379 Mouse PTP |
| C | 77 | 15.8 | 75.2 | 31236 | 12 | ADM74495 | Adm74495 Human car |
| C | 78 | 15.8 | 75.2 | 31718 | 4 | AAK90359 | Aak90359 Human dig |
| C | 79 | 15.8 | 75.2 | 31718 | 4 | AAK90360 | Aak90360 Human dig |
| C | 80 | 15.8 | 75.2 | 31718 | 4 | AAK73104 | Aak73104 Human imm |
| C | 81 | 15.8 | 75.2 | 31718 | 4 | AAK87573 | Aak87573 Human imm |
| C | 82 | 15.8 | 75.2 | 31718 | 4 | AAK73120 | Aak73120 Human imm |
| C | 83 | 15.8 | 75.2 | 31718 | 4 | AAK87442 | Aak87442 Human imm |
| C | 84 | 15.8 | 75.2 | 31718 | 4 | AAK87443 | Aak87443 Human imm |
| C | 85 | 15.8 | 75.2 | 31718 | 4 | AAK87592 | Aak87592 Human imm |
| C | 86 | 15.8 | 75.2 | 31718 | 4 | AAK06415 | Aal06415 Human rep |
| C | 87 | 15.8 | 75.2 | 31718 | 4 | AAAL06416 | Aal06416 Human rep |
| C | 88 | 15.8 | 75.2 | 31718 | 5 | AAS39916 | Aas39916 Genomic s |
| C | 89 | 15.8 | 75.2 | 31718 | 5 | AAS39915 | Aas39915 Genomic s |
| C | 90 | 15.8 | 75.2 | 31718 | 9 | ADB32875 | AdB32875 Human nov |
| C | 91 | 15.8 | 75.2 | 31718 | 9 | ADB32876 | AdB32876 Human nov |
| C | 92 | 15.8 | 75.2 | 31718 | 12 | ADN41665 | Adn41665 Novel hum |
| C | 93 | 15.8 | 75.2 | 31718 | 12 | ADN41666 | Adn41666 Novel hum |

c 94 15.8 75.2 63609 12 ADQ97537 Adq97537 Human can
95 15.8 75.2 110000 9 ADL12064.07 Continuation (8 of
96 15.8 75.2 122923 11 ACN44026 ACN44026 Human gen
c 97 15.8 75.2 170170 10 ADL13643 ADL13643 Osteoarthritis
98 15.4 73.3 20 6 AAD34672 AAD34672 DST CHS1
c 99 15.4 73.3 327 6 AAD34632 AAD34632 HBV infec
100 15.4 73.3 640 12 ADJ75740 ADJ75740 Marker ge

ALIGNMENTS

RESULT 1
ACC80536/c

ID ACC80536 standard; DNA; 21 BP.

XX

AC ACC80536;

XX

29-AUG-2003 (first entry)

XX

DE Exemplary sequence for method detecting HBV DNA in a sample.

XX

Hepatitis B virus; diagnosis; nuclease assay; ss;

KW

KW transcription-mediated amplification.

XX

OS Hepatitis B virus.

XX

PN WO2003031934-A2.

XX

PD 17-APR-2003.

XX

09-OCT-2002; 2002WO-US032367.

XX

09-OCT-2001; 2001US-0328492P.

PR

29-MAR-2002; 2002US-0368823P.

PR

02-JUL-2002; 2002US-0393561P.

XX

(CHIR) CHIRON CORP.

PA

PI Shyamala V;

XX

WPI; 2003-403124/38.

DR

New isolated hepatitis B virus (HBV) capture oligonucleotides, useful for detecting HBV infection in a biological sample, or in capturing HBV nucleic acids.

PT

PT

XX

PS Disclosure; Page 22; 50pp; English.

XX

The invention relates to a novel method of detecting hepatitis B virus (HBV) infections in e.g. blood samples from donors, by capturing and amplifying conserved regions of the HBV genome using a transcription-mediated amplification (TMA) method as well as a 5' nuclease assay. This sequence represents an exemplary replacement sequence for a target sequence (ACC80535) used in an internal control for the method of the invention. The new method is very sensitive and is able to detect about 100 infectious units (IU) of HBV in a viremic sample

XX

SQ Sequence 21 BP; 5 A; 6 C; 5 G; 5 T; 0 U; 0 Other;

XX

Query Match 100.0%; Score 21; DB 8; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.5;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CAGTGACATGCAGGCTAGCT 21

DB 21 CAGTGACATGCAGGCTAGCT 1

XX

RESULT 2

ACC80538

ID ACC80538 standard; DNA; 21 BP.

XX

AC ACC80538;

AC

XX

29-AUG-2003 (first entry)

XX

Internal control primer #2 for hepatitis B virus DNA detection method.

DE

XX

Primer; PCR; ss; hepatitis B virus; diagnosis; nuclease assay;

KW

KW transcription-mediated amplification.

XX

OS Hepatitis B virus.

XX

PH Key

misc_difference 1

FT

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FT

KW Primer; PCR; ss; hepatitis B virus; diagnosis; nuclease assay;
 KW transcription-mediated amplification.

OS Hepatitis B virus.

XX Key Location/Qualifiers

XX misc_difference 1

FT /tag= a
 FT /note= "linked to 6-PAM"

FT misc_difference 21

FT /tag= b
 FT /note= "linked to TAMRA"

XX WO2003031934-A2.

XX 17-APR-2003.

XX 09-OCT-2002; 2002WO-US032367.

XX 09-OCT-2001; 2001US-0328492P.

PR 29-MAR-2002; 2002US-0368823P.

PR 02-JUL-2002; 2002US-0393561P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2003-403124/38.

XX New isolated hepatitis B virus (HBV) capture oligonucleotides, useful for
 FT detecting HBV infection in a biological sample, or in capturing HBV
 FT nucleic acids.

XX Claim 13; Fig 2; 50pp; English.

XX The invention relates to a novel method of detecting hepatitis B virus
 CC (HBV) infections in e.g. blood samples from donors, by capturing and
 CC amplifying conserved regions of the HBV genome using a transcription-
 CC mediated amplification (TMA) method as well as a 5' nuclease assay. The
 CC method may also use an internal control sequence such as ACC80539, to
 CC determine the level of amplification and detection by the primers and
 CC probes used in the method of the invention. This sequence represents a
 CC primer used to amplify the internal control region DNA sequence. The new
 CC method is very sensitive and is able to detect about 100 infectious units
 CC (IU) of HBV in a viremic sample

XX Sequence 21 BP; 5 A; 6 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 8; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.5;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTTAGCT 21

Db 21 CAGTGACATGCAGGCTTAGCT 1

RESULT 4

ABZ59632/C

ID ABZ59632 standard; DNA; 21 BP.

XX

AC ABZ59632;

XX

DT 22-APR-2003 (first entry)

XX

DE Parvovirus B19 related internal control oligonucleotide SEQ ID NO:90.

XX

KW Human parvovirus B19; parvovirus B19; infection; virus; blood; plasma;

XX PCR primer; ss.

XX

OS B19 virus.

OS Synthetic.

XX

PN WO2003002753-A2.

XX

PD 09-JAN-2003.

XX

XX 28-JUN-2002; 2002WO-US020684.

XX

XX 28-JUN-2001; 2001US-0302077P.

PR 19-MAR-2002; 2002US-0365956P.

PR 29-MAR-2002; 2002US-0369224P.

XX

XX (CHIR) CHIRON CORP.

XX

XX Pichuanes S, Shyamala V;

XX

XX WPI; 2003-201510/19.

XX

XX Detecting a human parvovirus B19 infection in a biological sample to
 PT prevent viral transmission, comprises reacting a parvovirus B19 nucleic
 PT acid with a primer complementary to the 3'-terminal portion of the RNA
 PT target sequence.

XX Claim 8; Page 29; 148pp; English.

XX The present invention describes a method for detecting a human parvovirus
 CC B19 infection in a biological sample. The method comprises reacting the
 CC isolated parvovirus B19 nucleic acid with a first oligonucleotide
 CC consisting of a first primer containing a complexing sequence
 CC sufficiently complementary to the 3'-terminal portion of the RNA target
 CC sequence to complex with. Also described: (1) amplifying a target
 CC parvovirus B19 nucleotide sequence; (2) a polynucleotide comprising one
 CC of 47 700 base pair sequences (see ABZ59549 to ABZ59569, and ABZ59604 to
 CC ABZ59629); (3) a polynucleotide comprising either of 2 4678 base pair
 CC sequences (see ABZ59570 and ABZ59571); (4) an oligonucleotide primer
 CC consisting of a promoter region recognised by a DNA-dependent RNA
 CC polymerase operably linked to a human parvovirus B19-specific complexing
 CC sequence of 10-75 nucleotides; (5) an oligonucleotide probe comprising a
 CC parvovirus B19-specific hybridising sequence of 10-50 nucleotides linked
 CC to an acridinium ester label; and (6) a diagnostic test kit comprising an
 CC oligonucleotide primer of (4), and instructions for conducting the
 CC diagnostic test. The method is useful for detecting parvovirus infection
 CC in a biological sample, such as in blood products, to prevent
 CC transmission of the virus through blood and plasma derivatives or by
 CC close personal contact. ABZ59549 to ABZ59634 and ABP57262 to ABP57267
 CC represent sequences used in the exemplification of the present invention

XX Sequence 21 BP; 5 A; 6 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 9; Length 21;

Best Local Similarity 100.0%; Pred. No. 2.5;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CAGTGACATGCAGGCTTAGCT 21

Db 21 CAGTGACATGCAGGCTTAGCT 1

RESULT 5

ADIS3796

ID ADIS3796 standard; DNA; 21 BP.

XX

AC ADIS3796;

XX

DT 06-MAY-2004 (first entry)

XX

DE HAV internal control specific detection probe.

XX

KW HAV; nucleic acid amplification; nucleic acid detection; ss; probe.

XX

OS Hepatitis A virus.

OS Synthetic.

XX

XX Key Location/Qualifiers

XX modified_base 1

```

FT      /*tag= a
FT      /mod_base= 5'-TET labelled
FT      21
FT      /*tag= b
FT      /mod_base= 3'-TAMRA labelled
XX      WO2003106641-A2.
XX      24-DEC-2003.
XX
XX      12-JUN-2003; 2003WO-US018827.
XX
XX      12-JUN-2002; 2002US-0388544P.
XX      (CHIR ) CHIRON CORP.
XX      Shyamala V;
XX      WPI; 2004-082181/08.
XX      Hepatitis A virus specific primers and probes derived from conserved
XX      regions of the hepatitis A virus genome, useful in nucleic acid-based
XX      diagnostic tests for the detection of Hepatitis A virus in biological
XX      samples.
XX      Example 2; SEQ ID NO 18; 44pp; English.
XX
XX      The invention relates to Hepatitis A virus (HAV) specific primers and
XX      probes derived from conserved regions of the hepatitis A virus genome.
XX      The HAV-specific primers and probes are used in a method for detecting
XX      HAV in a biological sample. Also provided are capture oligonucleotides
XX      (Seq ID. No. 10)-(Seq ID. No. 14) which are used in a method for
XX      detecting HAV infection in a biological sample. The present sequence
XX      represents a detection probe specific for an internal control sequence.
XX      Sequence 21 BP; 5 A; 5 C; 6 G; 5 T; 0 U; 0 Other;
XX
XX      Query Match      100.0%; Score 21; DB 12; Length 21;
XX      Best Local Similarity 100.0%; Pred. No. 2.5;
XX      Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX      QY      1 CAGTGACATGCAGGTCTAGCT 21
XX      |||||
XX      Db      1 CAGTGACATGCAGGTCTAGCT 21
XX
XX      RESULT 6
XX      ADI53797
XX      ID      ADI53797 standard; DNA; 23 BP.
XX      AC      ADI53797;
XX      XX
XX      DT      06-MAY-2004 (first entry)
XX      XX
XX      DE      HAV internal control specific detection probe.
XX      XX
XX      KW      HAV; nucleic acid amplification; nucleic acid detection; ss; probe.
XX      XX
XX      OS      Hepatitis A virus.
XX      OS      Synthetic.
XX      XX
XX      FH      Key      Location/Qualifiers
XX      FT      modified_base 1
XX      FT      /*tag= a
XX      FT      /mod_base= 5'-TET labelled
XX      FT      23
XX      FT      modified_base
XX      FT      /*tag= b
XX      FT      /mod_base= 3'-TAMRA labelled
XX      XX
XX      PN      WO2003106641-A2.
XX      XX
XX      PD      24-DEC-2003.
XX
XX      PF      12-JUN-2003; 2003WO-US018827.
XX      XX
XX      PR      12-JUN-2002; 2002US-0388544P.
XX      XX      (CHIR ) CHIRON CORP.
XX      XX      Shyamala V;
XX      XX      WPI; 2004-082181/08.
XX      XX      Hepatitis A virus specific primers and probes derived from conserved
XX      regions of the hepatitis A virus genome, useful in nucleic acid-based
XX      diagnostic tests for the detection of Hepatitis A virus in biological
XX      samples.
XX      Example 2; SEQ ID NO 19; 44pp; English.
XX
XX      The invention relates to Hepatitis A virus (HAV) specific primers and
XX      probes derived from conserved regions of the hepatitis A virus genome.
XX      The HAV-specific primers and probes are used in a method for detecting
XX      HAV in a biological sample. Also provided are capture oligonucleotides
XX      (Seq ID. No. 10)-(Seq ID. No. 14) which are used in a method for
XX      detecting HAV infection in a biological sample. The present sequence
XX      represents a detection probe specific for an internal control sequence.
XX      Sequence 23 BP; 5 A; 7 C; 6 G; 5 T; 0 U; 0 Other;
XX
XX      Query Match      100.0%; Score 21; DB 12; Length 23;
XX      Best Local Similarity 100.0%; Pred. No. 2.5;
XX      Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX      QY      1 CAGTGACATGCAGGTCTAGCT 21
XX      |||||
XX      Db      3 CAGTGACATGCAGGTCTAGCT 23
XX
XX      RESULT 7
XX      ADQ30670
XX      ID      ADQ30670 standard; DNA; 23 BP.
XX      XX      ADQ30670;
XX      AC      ADQ30670;
XX      XX
XX      DT      23-SEP-2004 (first entry)
XX      XX
XX      DE      West Nile Virus internal control probe #1.
XX      XX
XX      KW      ss; primer; West Nile Virus; diagnosis.
XX      XX
XX      OS      West Nile virus.
XX      XX      WO2004055159-A2.
XX      PN      WO2004055159-A2.
XX      XX
XX      PD      01-JUL-2004.
XX      XX
XX      PF      05-DEC-2003; 2003WO-US038750.
XX      XX
XX      PR      12-DEC-2002; 2002US-0432850P.
XX      PR      20-JUN-2003; 2003US-0480431P.
XX      XX      (CHIR ) CHIRON CORP.
XX      PA      (CHIR ) CHIRON CORP.
XX      XX      Shyamala V;
XX      PI      Shyamala V;
XX      XX      WPI; 2004-488058/46.
XX      XX      New isolated oligonucleotides for accurately diagnosing West Nile virus
XX      infection or for capturing, detecting and quantitating West Nile virus in
XX      blood samples.
XX      Claim 29; SEQ ID NO 40; 56pp; English.
XX
XX      The invention relates to an isolated oligonucleotide not more than 60
XX      nucleotides in length comprising a nucleotide sequence (S1) of at least

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CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to a probe to the internal control sequence for the detection
 CC of WNV sequences using the oligonucleotides of the invention.

XX Sequence 23 BP; 5 A; 5 C; 6 G; 5 T; 0 U; 2 Other;

Query Match 100.0%; Score 21; DB 12; Length 23;
 Best Local Similarity 100.0%; Pred. No. 2.5; Mismatches 0; Indels 0; Gaps 0;
 Matches 21; Conservative 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21
 |||||
 Db 2 CAGTGACATGCAGGCTTAGCT 22

RESULT 8
 ADQ30671
 ID ADQ30671 standard; DNA; 25 BP.

AC ADQ30671;

XX 23-SEP-2004 (first entry)

XX West Nile Virus internal control probe #2.

XX ss; primer; West Nile Virus; diagnosis.

XX West Nile virus.

XX WO2004055159-A2.

XX 01-JUL-2004.

XX 05-DEC-2003; 2003WO-US038750.

XX 12-DEC-2002; 2002US-0432850P.

XX 20-JUN-2003; 2003US-0480431P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2004-488058/46.

XX New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.

XX Claim 29; SEQ ID NO 41; 56pp; English.

XX The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence

CC corresponds to a probe to the internal control sequence for the detection
 CC of WNV sequences using the oligonucleotides of the invention.

XX Sequence 25 BP; 5 A; 7 C; 6 G; 5 T; 0 U; 2 Other;

Query Match 100.0%; Score 21; DB 12; Length 25;
 Best Local Similarity 100.0%; Pred. No. 2.5; Mismatches 0; Indels 0; Gaps 0;
 Matches 21; Conservative 0;

QY 1 CAGTGACATGCAGGCTTAGCT 21
 |||||
 Db 4 CAGTGACATGCAGGCTTAGCT 24

RESULT 9
 ABZ59634

ID ABZ59634 standard; DNA; 681 BP.

XX AC ABZ59634;

XX 22-APR-2003 (first entry)

XX Exemplary internal control nucleotide sequence SEQ ID NO:92.

XX Human parvovirus B19; parvovirus B19; infection; virus; blood; plasma;
 XX gene; ds.

XX Synthetic.

XX WO2003002753-A2.

XX 09-JAN-2003.

XX 28-JUN-2002; 2002WO-US020684.

XX 28-JUN-2001; 2001US-0302077P.

XX 19-MAR-2002; 2002US-0365956P.

XX 29-MAR-2002; 2002US-0369224P.

XX (CHIR) CHIRON CORP.

XX Pichuanes S, Shyamala V;

XX WPI; 2003-201510/19.

XX Detecting a human parvovirus B19 infection in a biological sample to
 PT prevent viral transmission, comprises reacting a parvovirus B19 nucleic
 PT acid with a primer complementary to the 3'-terminal portion of the RNA
 PT target sequence.

XX Claim 7; Fig 12; 148pp; English.

XX The present invention describes a method for detecting a human parvovirus
 CC B19 infection in a biological sample. The method comprises reacting the
 CC isolated parvovirus B19 nucleic acid with a first oligonucleotide
 CC consisting of a first primer containing a complexing sequence
 CC sufficiently complementary to the 3'-terminal portion of the RNA target
 CC sequence to complex with. Also described: (1) amplifying a target
 CC parvovirus B19 nucleotide sequence; (2) a polynucleotide comprising one
 CC of 47 700 base pair sequences (see ABZ59549 to ABZ59569, and ABZ59604 to
 CC ABZ59629); (3) a polynucleotide comprising either of 2 4678 base pair
 CC sequences (see ABZ59570 and ABZ59571); (4) an oligonucleotide primer
 CC consisting of a promoter region recognised by a DNA-dependent RNA
 CC polymerase operably linked to a human parvovirus B19-specific complexing
 CC sequence of 10-75 nucleotides; (5) an oligonucleotide probe comprising a
 CC parvovirus B19-specific hybridising sequence of 10-50 nucleotides linked
 CC to an acridinium ester label; and (6) a diagnostic test kit comprising an
 CC oligonucleotide primer of (4), and instructions for conducting the
 CC diagnostic test. The method is useful for detecting parvovirus infection
 CC in a biological sample, such as in blood products, to prevent
 CC transmission of the virus through blood and plasma derivatives or by
 CC close personal contact. ABZ59549 to ABZ59634 and ABP57262 to ABP57267
 CC represent sequences used in the exemplification of the present invention

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XX SQ Sequence 681 BP; 206 A; 138 C; 137 G; 200 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 21; DB 9; Length 681;
XX Best Local Similarity 100.0%; Pred. No. 3.4;
XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 CAGTGACATGCAGGCTTAGCT 21
DQ 62 CAGTGACATGCAGGCTTAGCT 82
XX
RESULT 10
AD153795
ID AD153795 standard; DNA; 727 BP.
AC AD153795;
XX
DT 06-MAY-2004 (first entry)
XX
DE HAV internal control sequence.
XX
KW HAV; nucleic acid amplification; nucleic acid detection; ds.
XX
OS Hepatitis A virus.
OS Synthetic.
XX
FN WO2003106641-A2.
XX
PD 24-DEC-2003.
XX
PF 12-JUN-2003; 2003WO-US018827.
XX
PR 12-JUN-2002; 2002US-0388544P.
XX
PA (CHIR ) CHIRON CORP.
XX
PI Shyamala V;
XX
WPI; 2004-082181/08.
XX
Hepatitis A virus specific primers and probes derived from conserved
PT regions of the hepatitis A virus genome, useful in nucleic acid-based
PT diagnostic tests for the detection of Hepatitis A virus in biological
PT samples.
XX
XX Example 2; SEQ ID NO 17; 44pp; English.
XX
CC The invention relates to Hepatitis A virus (HAV) specific primers and
CC probes derived from conserved regions of the hepatitis A virus genome.
CC The HAV-specific primers and probes are used in a method for detecting
CC HAV in a biological sample. Also provided are capture oligonucleotides
CC (Seq ID. No. 10)-(Seq ID. No. 14) which are used in a method for
CC detecting HAV infection in a biological sample. The present sequence
CC represents an internal control sequence used as a control for target
CC capture and amplification.
XX
XX Sequence 727 BP; 147 A; 169 C; 186 G; 225 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 21; DB 12; Length 727;
XX Best Local Similarity 100.0%; Pred. No. 3.4;
XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 CAGTGACATGCAGGCTTAGCT 21
DQ 581 CAGTGACATGCAGGCTTAGCT 601
XX
RESULT 11
ADQ30647
ID ADQ30647 standard; DNA; 967 BP.
XX
AC ADQ30647;
XX
XX Sequence 967 BP; 273 A; 206 C; 272 G; 216 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 21; DB 12; Length 967;
XX Best Local Similarity 100.0%; Pred. No. 3.5;
XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 CAGTGACATGCAGGCTTAGCT 21
DQ 153 CAGTGACATGCAGGCTTAGCT 173
XX
RESULT 12
ACC80539
ID ACC80539 standard; DNA; 1696 BP.
XX
AC ACC80539;
XX
DT 29-AUG-2003 (first entry)
XX
DE Internal control region for hepatitis B virus DNA detection method.
XX
KW Hepatitis B virus; diagnosis; nucleic acid assay; internal control region;
KW transcription-mediated amplification; ds.
XX
OS Hepatitis B virus.
XX
FN WO2003031934-A2.
XX
XX Sequence 967 BP; 273 A; 206 C; 272 G; 216 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 21; DB 12; Length 967;
XX Best Local Similarity 100.0%; Pred. No. 3.5;
XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 CAGTGACATGCAGGCTTAGCT 21
DQ 153 CAGTGACATGCAGGCTTAGCT 173
XX
RESULT 12
ACC80539
ID ACC80539 standard; DNA; 1696 BP.
XX
AC ACC80539;
XX
DT 29-AUG-2003 (first entry)
XX
DE Internal control region for hepatitis B virus DNA detection method.
XX
KW Hepatitis B virus; diagnosis; nucleic acid assay; internal control region;
KW transcription-mediated amplification; ds.
XX
OS Hepatitis B virus.
XX
FN WO2003031934-A2.
XX

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XX PD 17-APR-2003.
 XX PF 09-OCT-2002; 2002WO-US032367.
 XX PR 09-OCT-2001; 2001US-0328492P.
 PR 29-MAR-2002; 2002US-0368823P.
 XX 02-JUL-2002; 2002US-0393561P.
 XX PA (CHIR) CHIRON CORP.
 XX PI Shyamala V;
 XX WPI; 2003-403124/38.
 XX New isolated hepatitis B virus (HBV) capture oligonucleotides, useful for
 PT detecting HBV infection in a biological sample, or in capturing HBV
 PT nucleic acids.
 XX Disclosure; Fig 3; 50pp; English.
 XX The invention relates to a novel method of detecting hepatitis B virus
 CC (HBV) infections in e.g. blood samples from donors, by capturing and
 CC amplifying conserved regions of the HBV genome using a transcription-
 CC mediated amplification (TMA) method as well as a 5' nuclease assay.
 CC method may also use an internal control sequence (this sequence), to
 CC determine the level of amplification and detection by the primers and
 CC probes used in the method of the invention. The new method is very
 CC sensitive and is able to detect about 100 infectious units (IU) of HBV in
 CC a viresmic sample
 XX Sequence 1696 BP; 359 A; 462 C; 386 G; 489 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 21; DB 8; Length 1696;
 Best Local Similarity 100.0%; Pred. No. 3.7;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CAGTGACATGCAGGCTCTAGCT 21
 Db 1428 CAGTGACATGCAGGCTCTAGCT 1448
 RESULT 13
 ADE71204/C
 ID ADE71204 standard; DNA; 5446 BP.
 XX AC ADE71204;
 XX 29-JAN-2004 (first entry)
 XX Novel human protein coding sequence #20.
 XX human; novel protein; drug; gene; ds.
 XX Homo sapiens.
 XX JP2002345493-A.
 XX 03-DEC-2002.
 XX 29-MAR-2001; 2002JP-00049046.
 XX 29-MAR-2001; 2001JP-00095524.
 XX (KAZU-) ZH KAZUSA DNA KENKYUSHO.
 XX WPI; 2003-460885/44.
 DR P-PSDB; ADE71266.
 XX A gene and a protein encoded by it, used in drugs.
 XX Claim 1; SEQ ID NO 20; 257pp; Japanese.

CC The invention comprises the amino acid and coding sequences of novel
 CC human proteins. The DNA and protein sequences of the invention are used
 CC in drugs. The present DNA sequence encodes a novel human protein of the
 CC invention.
 XX Sequence 5446 BP; 1477 A; 1320 C; 1292 G; 1357 T; 0 U; 0 Other;
 SQ Query Match 82.9%; Score 17.4; DB 10; Length 5446;
 Best Local Similarity 94.7%; Pred. No. 2.3e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 CAGTGACATGCAGGCTCTAG 19
 Db 833 CAGTGACAGGAGGCTCTAG 815
 RESULT 14
 ABZ73855/C
 ID ABZ73855 standard; DNA; 28482 BP.
 XX AC ABZ73855;
 XX 12-MAY-2003 (first entry)
 XX Secreted protein gene 64 genomic fragment HCEGX05, SEQ ID NO:1002.
 DE Human; secreted protein; cancer; tumour; hyperproliferative disorder;
 XX autoimmune disorder; inflammation; angiogenic diseases; AIDS;
 XX acquired immunodeficiency syndrome; hepatitis; anaemia; wound healing;
 KW drug screening; chromosome identification; chromosome mapping;
 KW cytostatic; gene therapy; antiinflammatory; immunomodulator; anti-HIV;
 KW antianaemic; vulnery; chromosome 20q13; gene; ds.
 XX Homo sapiens.
 XX W0200277013-A2.
 XX 03-OCT-2002.
 XX 26-MAR-2002; 2002WO-US009370.
 XX 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX Rosen CA, Ruben SM;
 WPI; 2003-040578/03.
 XX New human secreted proteins and nucleic acids, useful for detecting or
 PT treating cancer or other hyperproliferative disorders, autoimmune
 PT disorders, inflammatory disorders, HIV disease, hepatitis or anemia.
 XX Disclosure; Page 1651-1658; 2474pp; English.
 XX ABZ73281-ABZ73697 represent cDNAs corresponding to 391 human secreted
 CC protein genes, and ABP00947-ABP01363 represent the proteins they encode.
 CC ABZ73698-ABZ74687 represent human secreted protein genomic fragments. The
 CC invention also encompasses antibodies specific for the secreted proteins,
 CC the use of the secreted proteins in drug screening and recombinant
 CC vectors and host cells comprising a nucleic acid of the invention. The
 CC secreted proteins are thought to be involved in biological activities
 CC associated with cellular signalling, cellular differentiation, cell
 CC migration, prohormone activation and neurotransmitter activity. The
 CC secreted proteins, nucleic acids encoding them, antibodies or antibody
 CC fragments specific for the secreted proteins, and modulators of protein
 CC activity are useful for diagnosing or treating cancers or other
 CC hyperproliferative disorders. Additionally, the secreted proteins and
 CC their nucleic acids may also be used in the treatment of autoimmune
 CC disorders, inflammatory disorders, diseases involving angiogenesis, AIDS
 CC (acquired immunodeficiency syndrome), hepatitis, anaemia, and to promote

CC wound healing. Nucleic acids of the invention may be used for chromosome
CC identification, chromosome mapping, in gene therapy, for identifying
CC individuals from minute biological samples, as hybridisation probes, and
CC as molecular weight markers. The present sequence represents a human
CC secreted protein genomic fragment referred to in the disclosure of the
CC invention

XX Sequence 28482 BP; 7636 A; 6245 C; 6763 G; 7838 T; 0 U; 0 Other;

SQ Query Match 82.9%; Score 17.4; DB 8; Length 28482;

Best Local Similarity 94.7%; Pred. No. 2.6e+02;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CAGTGACATGCAGGCTCTAG 19

|||||||

DB 12331 CAGTGACAGGCAGGCTCTAG 12313

RESULT 15

ADA44262/c

ID ADA44262 standard; DNA; 28482 BP.

XX AC ADA44262;

XX DT 20-NOV-2003 (first entry)

XX DE Human secreted protein DNA SEQ ID 455.

XX KW Gene therapy; human; Antidiabetic; Anorectic; Ophthalmological;

XX KW Neuroprotective; Cerebroprotective; Antianemic; ds.

XX OS Homo sapiens.

XX PN WO2003000865-A2.

XX PD 03-JAN-2003.

XX PF 26-MAR-2002; 2002WO-US009105.

XX PR 27-MAR-2001; 2001US-0278650P.

XX PR 12-SEP-2001; 2001US-00950082.

XX PR 12-SEP-2001; 2001US-00950083.

XX PA (HUMA-) HUMAN GENOME SCI INC.

XX PI Rosen CA, Ruben SM;

XX DR WPI; 2003-184045/18.

XX PT A human secreted protein and nucleic acids useful for preparing a
PT diagnostic or pharmaceutical composition for diagnosing or treating
PT diabetes or conditions related to diabetes, e.g. hyperglycemia, obesity,
PT retinopathy, neuropathy.

XX PS Disclosure; SEQ ID NO 455; 701pp; English.

XX CC The invention relates to novel genes and their fragments which are useful
CC for preventing, treating or ameliorating medical conditions e.g. by
CC protein or gene therapy. The genes are isolated from a range of human
CC tissues disclosed in the specification. The nucleic acids and proteins
CC are useful in the diagnosis, treatment and prevention of conditions
CC related to diabetes, e.g. hyperglycaemia, obesity, retinopathy,
CC polynuropathy, atherosclerosis, anaemia, stroke, gangrene, impotence,
CC infection, cataract, renal disorders, or endocrine disorders. The present
CC sequence was used to illustrate the invention.

XX SQ Sequence 28482 BP; 7636 A; 6245 C; 6763 G; 7838 T; 0 U; 0 Other;

Query Match

Best Local Similarity 82.9%; Score 17.4; DB 8; Length 28482;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CAGTGACATGCAGGCTCTAG 19

DB 12331 CAGTGACAGGCAGGCTCTAG 12313

RESULT 16

ABZ74517/c

ID ABZ74517 standard; DNA; 32681 BP.

XX AC ABZ74517;

XX DT 12-MAY-2003 (first entry)

XX DE Secreted protein gene 329 genomic fragment HTLT80, SEQ ID NO:1664.

XX KW Human; secreted protein; cancer; tumour; hyperproliferative disorder;
XX KW autoimmune disorder; inflammation; angiogenic diseases; AIDS;
XX KW acquired immunodeficiency syndrome; hepatitis; anaemia; wound healing;
XX KW drug screening; chromosome identification; chromosome mapping;
XX KW cytostatic; gene therapy; antiinflammatory; immunomodulator; anti-HIV;
XX KW antianaemic; vulnery; chromosome 20q11.21-13.11; gene; ds.

XX OS Homo sapiens.

XX PN WO200277013-A2.

XX PD 03-OCT-2002.

XX PF 26-MAR-2002; 2002WO-US009370.

XX PR 27-MAR-2001; 2001US-0278650P.

XX PR 12-SEP-2001; 2001US-00950082.

XX PR 12-SEP-2001; 2001US-00950083.

XX PA (HUMA-) HUMAN GENOME SCI INC.

XX PI Rosen CA, Ruben SM;

XX DR WPI; 2003-040578/03.

XX PT New human secreted proteins and nucleic acids, useful for detecting or
PT treating cancer or other hyperproliferative disorders, autoimmune
PT disorders, inflammatory disorders, HIV disease, hepatitis or anaemia.

XX PS Disclosure; Page 2245-2253; 2474pp; English.

XX CC ABZ73281-ABZ73697 represent cDNAs corresponding to 391 human secreted
CC protein genes, and ABP00947-ABP01363 represent the proteins they encode.
CC ABZ73698-ABZ74687 represent human secreted protein genomic fragments. The
CC invention also encompasses antibodies specific for the secreted proteins,
CC the use of the secreted proteins in drug screening and recombinant
CC vectors and host cells comprising a nucleic acid of the invention. The
CC secreted proteins are thought to be involved in biological activities
CC associated with cellular signalling, cellular differentiation, cell
CC migration, prohormone activation and neurotransmitter activity. The
CC fragments specific for the secreted proteins, and modulators of protein
CC activity are useful for diagnosing or treating cancers or other
CC hyperproliferative disorders. Additionally, the secreted proteins and
CC their nucleic acids may also be used in the treatment of autoimmune
CC disorders, inflammatory disorders, diseases involving angiogenesis, AIDS
CC (acquired immunodeficiency syndrome), hepatitis, anaemia, and to promote
CC wound healing. Nucleic acids of the invention may be used for chromosome
CC identification, chromosome mapping, in gene therapy, for identifying
CC individuals from minute biological samples, as hybridisation probes, and
CC as molecular weight markers. The present sequence represents a human
CC secreted protein genomic fragment referred to in the disclosure of the
CC invention

XX SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;

Query Match 82.9%; Score 17.4; DB 8; Length 32681;

Best Local Similarity 94.7%; Pred. No. 2.7e+02;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTTAG 19
 DB 16526 CAGTGACAGGCAGGTCTTAG 16508

RESULT 17
 AB273854/c
 ID AB273854 standard; DNA; 32681 BP.
 XX AC AB273854;
 XX DT 12-MAY-2003 (first entry)
 XX DE Secreted protein gene 64 genomic fragment HCEGX05, SEQ ID NO:1001.
 XX DE Human; secreted protein; cancer; tumour; hyperproliferative disorder;
 XX KW autoimmune disorder; inflammation; angiogenic diseases; AIDS;
 XX KW acquired immunodeficiency syndrome; hepatitis; anaemia; wound healing;
 XX KW drug screening; chromosome identification; chromosome mapping;
 XX KW cytostatic; gene therapy; antiinflammatory; immunomodulator; anti-HIV;
 XX KW antianaemic; vulnery; chromosome 20q13; gene; da.
 XX OS Homo sapiens.
 XX PN WO200277013-A2.
 XX PD 03-OCT-2002.
 XX PF 26-MAR-2002; 2002WO-US009370.
 XX PR 27-MAR-2001; 2001US-0278650P.
 XX PR 12-SEP-2001; 2001US-00950082.
 XX PR 12-SEP-2001; 2001US-00950083.
 XX PA (HUMA-) HUMAN GENOME SCI INC.
 XX PI Rosen CA, Ruben SM;
 XX WPI; 2003-040578/03.
 XX PT New human secreted proteins and nucleic acids, useful for detecting or
 XX PT treating cancer or other hyperproliferative disorders, autoimmune
 XX PT disorders, inflammatory disorders, HIV disease, hepatitis or anemia.
 XX PS Disclosure; Page 1643-1651; 2474pp; English.
 XX CC AB273281-AB273697 represent cDNAs corresponding to 391 human secreted
 CC protein genes, and ABP00947-ABP01363 represent the proteins they encode.
 CC CC AB273698-AB274687 represent human secreted protein genomic fragments. The
 CC invention also encompasses antibodies specific for the secreted proteins,
 CC the use of the secreted proteins in drug screening and recombinant
 CC vectors and host cells comprising a nucleic acid of the invention. The
 CC secreted proteins are thought to be involved in biological activities
 CC associated with cellular signalling, cellular differentiation, cell
 CC migration, pro-hormone activation and neurotransmitter activity. The
 CC secreted proteins, nucleic acids encoding them, antibodies or antibody
 CC fragments specific for the secreted proteins, and modulators of protein
 CC activity are useful for diagnosing or treating cancers or other
 CC hyperproliferative disorders. Additionally, the secreted proteins and
 CC their nucleic acids may also be used in the treatment of autoimmune
 CC disorders, inflammatory disorders, diseases involving angiogenesis, AIDS
 CC (acquired immunodeficiency syndrome), hepatitis, anaemia, and to promote
 CC wound healing. Nucleic acids of the invention may be used for chromosome
 CC identification, chromosome mapping, in gene therapy, for identifying
 CC individuals from minute biological samples, as hybridisation probes, and
 CC as molecular weight markers. The present sequence represents a human
 CC secreted protein genomic fragment referred to in the disclosure of the
 CC invention
 XX SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 8; Length 32681;
 Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTTAG 19
 DB 16526 CAGTGACAGGCAGGTCTTAG 16508

RESULT 18
 ADA98915/c
 ID ADA98915 standard; DNA; 32681 BP.
 XX AC ADA98915;
 XX DT 20-NOV-2003 (first entry)
 XX DE Human secreted protein-related DNA sequence #508.
 XX DE human; secreted protein; cardiovascular disorder; arrhythmia;
 XX KW atherosclerosis; stroke; endocarditis; congestive heart failure;
 XX KW rheumatic heart disease; cardiomyopathy; hemorrhoids; varicose veins;
 XX KW migraine; thrombosis; neural disorder; immune system disorder;
 XX KW muscular disorder; reproductive disorder; gastrointestinal disorder;
 XX KW pulmonary disorder; renal disorder; proliferative disorder; cancer; da.
 XX OS Homo sapiens.
 XX PN WO2003004623-A2.
 XX PD 16-JAN-2003.
 XX PF 26-MAR-2002; 2002WO-US009922.
 XX PR 27-MAR-2001; 2001US-0278650P.
 XX PR 12-SEP-2001; 2001US-00950082.
 XX PR 12-SEP-2001; 2001US-00950083.
 XX PA (HUMA-) HUMAN GENOME SCI INC.
 XX PI Rosen CA, Ruben SM;
 XX WPI; 2003-247946/24.
 XX PT New human secreted polypeptide and nucleic acid molecules, useful for
 XX PT diagnosing, preventing, prognosticating or treating cardiovascular
 XX PT disorders (e.g. arrhythmia, atherosclerosis, cardiomyopathy, or
 XX PT thrombosis).
 XX PS Disclosure; SEQ ID NO 1024; 1572pp; English.
 XX CC The invention comprises the amino acid and coding sequence of human
 CC secreted proteins. The DNA and protein sequences of the invention are
 CC useful in the treatment of cardiovascular disorders, such as: arrhythmia,
 CC atherosclerosis, stroke, endocarditis, congestive heart failure,
 CC rheumatic heart disease, cardiomyopathy, hemorrhoids, varicose veins,
 CC migraine, or thrombosis. The DNA and protein sequences may also be used
 CC for treating or preventing: neural disorders, immune system disorders,
 CC muscular disorders, reproductive disorders, gastrointestinal disorders,
 CC pulmonary disorders, renal disorders, proliferative disorders and/or
 CC cancerous diseases. The present DNA sequence is used in the
 CC exemplification of the invention. NOTE: The present sequence is shown on
 CC the WIPO website.
 XX SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 8; Length 32681;
 Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAGTGACATGCAGGTCTTAG 19
 DB 16526 CAGTGACAGGCAGGTCTTAG 16508

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RESULT 19
ADA44261/c
ID ADA44261 standard; DNA; 32681 BP.
XX
XX AC ADA44261;
XX
XX DT 20-NOV-2003 (first entry)
XX
XX DE Human secreted protein DNA SEQ ID 454.
XX
XX KW Gene therapy; human; Antidiabetic; Anorectic; Ophthalmological;
XX Neuroprotective; Cerebroprotective; Antianemic; ds.
XX
XX OS Homo sapiens.
XX
XX PN WO2003000865-A2.
XX
XX PD 03-JAN-2003.
XX
XX PF 26-MAR-2002; 2002WO-US009105.
XX
XX PR 27-MAR-2001; 2001US-0278650P.
XX 12-SEP-2001; 2001US-00950082.
XX 12-SEP-2001; 2001US-00950083.
XX
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX PI Rosen CA, Ruben SM;
XX
XX PF WI; 2003-184045/18.
XX
XX PT A human secreted protein and nucleic acids useful for preparing a
diagnostic or pharmaceutical composition for diagnosing or treating
diabetes or conditions related to diabetes, e.g. hyperglycemia, obesity,
retinopathy, neuropathy.
XX
XX PS Disclosure; SEQ ID NO 454; 701pp; English.
XX
XX CC The invention relates to novel genes and their fragments which are useful
for preventing, treating or ameliorating medical conditions e.g. by
protein or gene therapy. The genes are isolated from a range of human
tissues disclosed in the specification. The nucleic acids and proteins
are useful in the diagnosis, treatment and prevention of conditions
related to diabetes, e.g. hyperglycaemia, obesity, retinopathy,
polynuropathy, atherosclerosis, anaemia, stroke, gangrene, impotence,
infection, cataract, renal disorders, or endocrine disorders. The present
sequence was used to illustrate the invention.
XX
XX SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
Query Match 82.9%; Score 17.4; DB 8; Length 32681;
Best Local Similarity 94.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

XX QY 1 CAGTGACATGCAGGCTCTAG 19
||||| |||||||
DB 16526 CAGTGACAGGCAGGCTCTAG 16508

RESULT 20
ADA44519/c
ID ADA44519 standard; DNA; 32681 BP.
XX
XX AC ADA44519;
XX
XX DT 20-NOV-2003 (first entry)
XX
XX DE Human secreted protein DNA SEQ ID 712.
XX
XX KW Gene therapy; human; Antidiabetic; Anorectic; Ophthalmological;
XX Neuroprotective; Cerebroprotective; Antianemic; ds.
XX
XX OS Homo sapiens.
XX
XX PN WO2003000865-A2.
XX
XX PD 03-JAN-2003.
XX
XX PF 26-MAR-2002; 2002WO-US009105.
XX
XX PR 27-MAR-2001; 2001US-0278650P.
XX 12-SEP-2001; 2001US-00950082.
XX 12-SEP-2001; 2001US-00950083.
XX
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX PI Rosen CA, Ruben SM;
XX
XX PF WI; 2003-184045/18.
XX
XX PT A human secreted protein and nucleic acids useful for preparing a
diagnostic or pharmaceutical composition for diagnosing or treating
diabetes or conditions related to diabetes, e.g. hyperglycemia, obesity,
retinopathy, neuropathy.
XX
XX PS Disclosure; SEQ ID NO 454; 701pp; English.
XX
XX CC The invention relates to novel genes and their fragments which are useful
for preventing, treating or ameliorating medical conditions e.g. by
protein or gene therapy. The genes are isolated from a range of human
tissues disclosed in the specification. The nucleic acids and proteins
are useful in the diagnosis, treatment and prevention of conditions
related to diabetes, e.g. hyperglycaemia, obesity, retinopathy,
polynuropathy, atherosclerosis, anaemia, stroke, gangrene, impotence,
infection, cataract, renal disorders, or endocrine disorders. The present
sequence was used to illustrate the invention.
XX
XX SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
Query Match 82.9%; Score 17.4; DB 8; Length 32681;
Best Local Similarity 94.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

XX QY 1 CAGTGACATGCAGGCTCTAG 19
||||| |||||||
DB 16526 CAGTGACAGGCAGGCTCTAG 16508
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OS Homo sapiens.
XX
XX PN WO2003000865-A2.
XX
XX PD 03-JAN-2003.
XX
XX PF 26-MAR-2002; 2002WO-US009105.
XX
XX PR 27-MAR-2001; 2001US-0278650P.
XX 12-SEP-2001; 2001US-00950082.
XX 12-SEP-2001; 2001US-00950083.
XX
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX
XX PI Rosen CA, Ruben SM;
XX
XX PF WI; 2003-184045/18.
XX
XX PT A human secreted protein and nucleic acids useful for preparing a
diagnostic or pharmaceutical composition for diagnosing or treating
diabetes or conditions related to diabetes, e.g. hyperglycemia, obesity,
retinopathy, neuropathy.
XX
XX PS Disclosure; SEQ ID NO 712; 701pp; English.
XX
XX CC The invention relates to novel genes and their fragments which are useful
for preventing, treating or ameliorating medical conditions e.g. by
protein or gene therapy. The genes are isolated from a range of human
tissues disclosed in the specification. The nucleic acids and proteins
are useful in the diagnosis, treatment and prevention of conditions
related to diabetes, e.g. hyperglycaemia, obesity, retinopathy,
polynuropathy, atherosclerosis, anaemia, stroke, gangrene, impotence,
infection, cataract, renal disorders, or endocrine disorders. The present
sequence was used to illustrate the invention.
XX
XX SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
Query Match 82.9%; Score 17.4; DB 8; Length 32681;
Best Local Similarity 94.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

XX QY 1 CAGTGACATGCAGGCTCTAG 19
||||| |||||||
DB 16526 CAGTGACAGGCAGGCTCTAG 16508

RESULT 21
ADC20949/c
ID ADC20949 standard; DNA; 32681 BP.
XX
XX AC ADC20949;
XX
XX DT 18-DEC-2003 (first entry)
XX
XX DE Human secreted protein-related DNA sequence #367.
XX
XX KW gene therapy; human; secreted protein; haemopoietic disorder;
XX haematological disorder; anaemia; haemophilia; inflammatory disorder;
XX inflammatory bowel disease; Crohn's disease; neoplastic disease; cancer;
XX leukaemia; wound healing; epithelial cell proliferation disorder;
XX immune disorder; autoimmune disorder; asthmatic disorder;
XX cardiovascular disorder; atherosclerosis; myocarditis;
XX infectious disease; HIV; AIDS; endocrine disorder; diabetes;
XX gastrointestinal disorder; duodenal ulcer; gastroenteritis; gene; ds.
XX
XX OS Homo sapiens.
XX
XX PN WO200292787-A2.
XX
XX PD 21-NOV-2002.
XX
XX PF 26-MAR-2002; 2002WO-US009257.
XX
```

PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 XX Rosen CA, Ruben SM;
 XX WPI; 2003-129287/12.
 XX
 XX New human secreted proteins and nucleic acid molecules, useful for
 PT preparing a diagnostic or pharmaceutical composition for diagnosing,
 PT preventing or treating hematopoietic or hematologic disorders, e.g.
 PT anemia or hemophilia.
 XX
 XX Disclosure; SEQ ID NO 903; 1512pp; English.
 XX
 XX The invention comprises the amino acid and coding sequences of human
 CC secreted proteins. The DNA and protein sequences of the invention are
 CC useful for detecting, preventing, diagnosing, prognosticating, treating
 CC or ameliorating: hematopoietic or haematological disorders (e.g. anaemia
 CC and haemophilia); inflammatory disorders (e.g. inflammatory bowel disease
 CC and Crohn's disease); neoplastic disease (e.g. cancer and leukaemia);
 CC wound healing and disorders of epithelial cell proliferation; immune
 CC disorders (e.g. autoimmune disorders and asthmatic disorders);
 CC cardiovascular disorders (e.g. atherosclerosis and myocarditis);
 CC infectious disease (e.g. HIV/AIDS); endocrine disorders (e.g. diabetes);
 CC and gastrointestinal disorders (e.g. duodenal ulcers and
 CC gastroenteritis). The present DNA sequence was used in the
 CC exemplification of the invention.
 XX
 SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 10; Length 32681;
 Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 CAGTGACATGCAGGCTCTAG 19
 Db 16526 CAGTGACAGGCAGGCTCTAG 16508
 RESULT 22
 ABZ68053/c
 ID ABZ68053 standard; DNA; 32681 BP.
 XX
 XX AC ABZ68053;
 XX
 XX DT 26-MAR-2003 (first entry)
 XX
 XX DE Human secreted protein encoding genomic DNA SEQ ID NO 1576.
 XX
 XX Human; secreted protein; nontropic; neuroprotective; cytostatic;
 KW virucide; dermatological; immunosuppressive; antiinflammatory; anti-HIV;
 KW vulnary; antibacterial; antiparkinsonian; antisking; antianaemic;
 KW antiarthritic; cancer; antihypertensive; hepatotropic; cerebroprotective;
 KW antiinflammatory; antiallergic; antidiabetic; antitumor; anticonvulsant;
 KW antifungal; antiparasitic; cardiac; immune disorder; infection; vaccine;
 KW cardiovascular disorder; neurological disease; nephrotropic;
 KW gene therapy; gene; ds.
 XX
 XX OS Homo sapiens.
 XX
 XX WO20027186-A2.
 XX
 XX PD 03-OCT-2002.
 XX
 XX PF 26-MAR-2002; 2002WO-US009188.
 XX
 XX PR 27-MAR-2001; 2001US-0278650P.
 PR 12-SEP-2001; 2001US-00950082.
 PR 12-SEP-2001; 2001US-00950083.
 XX

PA (HUMA-) HUMAN GENOME SCI INC.
 XX Rosen CA, Ruben SM;
 XX WPI; 2003-040583/03.
 XX
 XX New human secreted proteins encoded by genes contained in cDNA clones
 PT (e.g. HGCAC19), useful for preventing, treating or diagnosing e.g. AIDS,
 PT multiple sclerosis, herpes virus, leukemia, tick-borne encephalitis or
 PT West Nile fever.
 XX
 XX Disclosure; Page 2201-2209; 2423pp; English.
 XX
 XX The invention relates to novel human genes (ABZ66891-ABZ68209) and the
 CC encoded secreted proteins (ABP9470-ABP99872) useful for preventing,
 CC treating or ameliorating medical conditions e.g. by protein or gene
 CC therapy. The genes are isolated from a range of human tissues disclosed
 CC in the specification. The nucleic acids, proteins, antibodies and
 CC (ant)agonists are useful in the diagnosis, treatment and prevention of:
 CC (a) cancer, e.g. breast and ovarian cancer and other cancers of the
 CC adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
 CC lung or urogenital; (b) immune disorders e.g. Addison's disease,
 CC allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
 CC diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
 CC arthritis and ulcerative colitis; (c) cardiovascular disorders such as
 CC myocardial ischaemia; (d) wound healing; (e) neurological diseases e.g.
 CC cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
 CC bacterial, fungal and parasitic infections
 XX
 SQ Sequence 32681 BP; 8783 A; 7103 C; 7721 G; 9074 T; 0 U; 0 Other;
 Query Match 82.9%; Score 17.4; DB 10; Length 32681;
 Best Local Similarity 94.7%; Pred. No. 2.7e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 CAGTGACATGCAGGCTCTAG 19
 Db 16526 CAGTGACAGGCAGGCTCTAG 16508
 RESULT 23
 ABX46275
 ID ABX46275 standard; cDNA; 320 BP.
 XX
 XX AC ABX46275;
 XX
 XX DT 21-FEB-2003 (first entry)
 XX
 XX DE Bovine EST associated with lactation/muscle/fat deposition #11440.
 XX
 XX KW Bovine; ss; EST; expressed sequence tag; lactation; LMFD;
 KW muscle deposition; fat deposition; genome mapping; gene identification;
 KW gene analysis; cattle breeding.
 XX
 XX OS Bos Taurus.
 XX
 XX US2002137139-A1.
 XX
 XX PD 26-SEP-2002.
 XX
 XX PF 24-SEP-2001; 2001US-00960352.
 XX
 XX PR 12-JAN-1999; 99US-0115707P.
 PR 11-JAN-2000; 2000US-00480902.
 PR
 XX (BYAT/) BYATT J C.
 PA (MATH/) MATHIALAGAN N.
 PA (TAON/) TAO N.
 PA (WARR/) WARREN W C.
 XX
 XX Byatt JC, Mathialagan N, Tao N, Warren WC;
 XX WPI; 2003-110599/10.
 DR

XX New nucleic acid associated with lactation, and muscle and fat
PT deposition, useful for genome mapping, gene identification and analysis,
PT cattle breeding, or for genetically improving cattle.
XX
XX Claim 2; SEQ ID NO 11440; 245bp; English.
XX
XX The invention relates to a purified nucleic acid molecule associated with
CC lactation or muscle and fat deposition (designated LMPD), derived from
CC cattle, and the LMPD nucleic acid can specifically hybridise to a second
CC nucleic acid molecule comprising any of 15112 nucleotide sequences,
CC appearing as ABX34836-ABX49947, or complements of them. Also included are
CC ; (1) a transformed cell having a nucleic acid comprising an LMPD nucleic
CC acid linked to a promoter and a 3' non-translated sequence that
CC functions in the cell to cause termination of transcription and addition
CC of polyadenylated ribonucleotides to a 3' end of the mRNA molecule; and
CC (2) determining a level or pattern of a molecule in a bovine cell or
CC tissue comprising: (a) incubating a marker nucleic acid (comprising any
CC of the 15112 nucleic acid sequences or its complement or fragment) with a
CC complementary nucleic acid molecule obtained from the bovine cell or
CC tissue, where hybridisation between the marker nucleic acid and the
CC complementary nucleic acid permits the detection of the molecule; and (b)
CC detecting the level or pattern of the complementary nucleic acid, where
CC the detection of the complementary nucleic acid is predictive of the
CC level or pattern of the molecule. The LMPD nucleic acid is used for
CC determining a level or pattern of a molecule in a bovine cell or tissue.
CC It is useful for genome mapping, gene identification and analysis, cattle
CC breeding, preparation of constructs for use in cattle gene expression, or
CC for genetically improving cattle. The present sequence is one of the
CC 15112 bovine LMPD EST (expressed sequence tag) nucleic acids. Note: The
CC present sequence was not shown in the specification but was obtained in
CC electronic format from the USPTO web site:
CC seqdata.uspto.gov/sequence.html?DocID=20020137139
XX
XX Sequence 320 BP; 78 A; 80 C; 97 G; 65 T; 0 U; 0 Other;

Query Match 80.0%; Score 16.8; DB 8; Length 320;
Best Local Similarity 90.0%; Pred. No. 3.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ACTGACATGCAGGCTTAGCT 21
Db 144 AATGACATGCAGGCTTACCT 163

RESULT 24
AAS00624
ID AAS00624 standard; DNA; 36221 BP.
XX
XX AAS00624;
XX
XX 07-SEP-2001 (first entry)
XX Human death-associated protein 6 (DAXX) gene.
XX
XX Death-associated protein 6; DAXX; polymorphism; haplotype pair; human;
KW immune disorder; autoimmune disease; population diversity; ds;
KW paternity testing; anthropological lineage; forensic application.
XX
XX Homo sapiens.
XX
XX Key Location/Qualifiers
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FT variation /*tag= b
FT variation /*tag= c
FT variation /*tag= d
FT variation /*tag= e
FT variation replace(27620,T)
FT variation /*tag= e
FT variation replace(27788,G)

FT variation /*tag= f
FT replace(27806,T)
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FT /*tag= r
FT replace(30752,T)
FT /*tag= s
FT replace(31916,T)
FT /*tag= t
XX WO200125245-A2.
XX 12-APR-2001.
XX
XX 05-OCT-2000; 2000WO-US027487.
XX
XX 06-OCT-1999; 99US-0157909P.
XX (GENA-) GENAISSANCE PHARM INC.
XX
XX Chew A, Choi JY, Denton RR, Nandabalan K, Stephens JC;
XX WPI; 2001-308220/32.
XX
XX New human death-associated protein 6 (DAXX) gene variants comprising 19
PT polymorphic sites useful in studying the effect of variation on the
PT biological activity of DAXX and in developing drugs targeting the
PT protein.
XX
XX Claim 1; Fig 1; 97pp; English.
XX
XX The sequence represents a DNA encoding human death-associated protein 6
CC (DAXX). This gene may comprise one or more polymorphisms at specific
CC nucleotide positions to form one of nineteen possible polymorphic
CC variants. Associations between a trait and a genotype or a haplotype of
CC the DAXX gene can be identified by comparing the frequency of the
CC genotype or haplotype in a population exhibiting the trait with that of a
CC reference population. A higher frequency in the trait population
CC indicates an association. Methods involving genotyping or haplotyping of
CC the DAXX gene of an individual can lead to prediction of haplotype pairs
CC for the DAXX gene of related individuals, and may be useful in studying
CC the expression and biological function of DAXX, as well as in developing
CC drugs targeting this protein. Polymorphic variants of DAXX are useful in
CC studying the effect of the variation on the biological activity of DAXX
CC as well as on the binding affinity of candidate drugs targeting DAXX for
CC the treatment of autoimmune diseases and other immune disorders.
CC Polymorphism is also useful for studying population diversity,
CC anthropological lineage, paternity testing, forensic applications, and
CC for identifying associations between the DAXX genetic variation and a
CC trait such as level of drug response or susceptibility to disease. DAXX
CC proteins may be used to measure binding affinities of one or more
CC candidate drugs targeting the DAXX protein
XX

SQ Sequence 36221 BP; 8897 A; 8473 C; 9437 G; 9414 T; 0 U; 0 Other;
Query Match 80.0%; Score 16.8; DB 4; Length 36221;
Best Local Similarity 90.0%; Pred. No. 5.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AGTGACATGCAGGCTCTAGCT 21
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Db 21390 AGTGACAGGCAGTCTTAGCT 21409

RESULT 25

ACN45158
ID ACN45158 standard; DNA; 72705 BP.

XX AC ACN45158;

XX DT 18-NOV-2004 (first entry)

XX DE Human genomic sequence hCG25130.

XX KW Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.

XX OS Homo sapiens.

XX PN WO2003073826-A2.

XX PD 12-SEP-2003.

XX PF 28-FEB-2003; 2003WO-US006235.

XX PR 01-MAR-2002; 2002US-00087192.

XX PA (SAGR-) SAGRES DISCOVERY.

XX PI Morris DW;

XX DR WPI; 2003-328604/31.

XX PT Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
comprises a nucleotide sequence.

XX PS Claim 1; SEQ ID NO 1966; Opp; English.

XX The present invention relates to novel DNA and protein sequences which
are associated with carcinomas. The sequences are useful for: (i) for
screening drug candidates; (ii) for screening of bioactive agent capable
of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
a bioactive agent capable of modulating the activity of CAP; (iv) for
evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
(x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
determining Carcinoma Associated (CA) gene copy number. In addition, the
CA genes are useful as DNA vaccines and the CAP are useful as markers of
carcinoma including lymphoma. The present sequence is one such CA coding
sequence. Note: This patent is an equivalent to basic patent
US2002182586A1, for which no sequence data was published

SQ Sequence 72705 BP; 18277 A; 18952 C; 18052 G; 17424 T; 0 U; 0 Other;
Query Match 78.1%; Score 16.4; DB 11; Length 72705;
Best Local Similarity 94.4%; Pred. No. 8.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 AGTGACATGCAGGCTCTAG 19
||||| ||||| ||||| |||||
Db 32065 AGTGACATGCAGGCTCTAG 32082

Search completed: September 6, 2005, 20:39:33
Job time : 194.656 secs

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OM nucleic - nucleic search, using sw model

Run on: September 9, 2005, 11:24:12 ; Search time 1697 Seconds
(without alignments)
542.516 Million cell updates/sec

Title: US-10-729-421-52
Perfect score: 19
Sequence: 1 cggatgcccgcgtgttg 19

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 1981570

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 150 summaries

Database :

GenEmbl.*
1: gb_ba.*
2: gb_htg.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_ets.*
12: gb_ey.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|-------------|--------------------|
| 1 | 12.6 | 66.3 | 24 | 6 AR430777 | AR430777 Sequence |
| C 2 | 12.6 | 66.3 | 54 | 6 AX456376 | AX456376 Sequence |
| C 3 | 12.6 | 66.3 | 60 | 6 AR300561 | AR300561 Sequence |
| C 4 | 12.2 | 64.2 | 18 | 12 AB068012 | AB068012 Synthetic |
| 5 | 12.2 | 64.2 | 24 | 6 AX289630 | AX289630 Sequence |
| 6 | 12.2 | 64.2 | 30 | 6 AR079209 | AR079209 Sequence |
| C 7 | 12.2 | 64.2 | 31 | 6 AX249387 | AX249387 Sequence |
| C 8 | 12.2 | 64.2 | 40 | 6 AR135209 | AR135209 Sequence |
| C 9 | 12.2 | 64.2 | 40 | 6 AR146705 | AR146705 Sequence |
| C 10 | 12.2 | 64.2 | 40 | 6 AR152276 | AR152276 Sequence |
| C 11 | 12.2 | 64.2 | 40 | 6 AR157814 | AR157814 Sequence |
| 12 | 12.2 | 64.2 | 50 | 6 CQ005341 | CQ005341 Sequence |
| C 13 | 12.2 | 64.2 | 55 | 6 AR523798 | AR523798 Sequence |
| 14 | 12 | 63.2 | 49 | 6 AX772391 | AX772391 Sequence |
| C 15 | 11.8 | 62.1 | 20 | 6 CQ840727 | CQ840727 Sequence |
| C 16 | 11.8 | 62.1 | 24 | 6 AX155207 | AX155207 Sequence |
| 17 | 11.8 | 62.1 | 48 | 6 A97649 | A97649 Sequence 16 |
| 18 | 11.8 | 62.1 | 48 | 6 AR428977 | AR428977 Sequence |
| 19 | 11.8 | 62.1 | 48 | 6 BD081713 | BD081713 Peptide. |

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| AX165870 | Sequence | 50 | 6 | AX165870 | Sequence |
| A97650 | Sequence | 53 | 6 | A97650 | Sequence |
| AR428978 | Sequence | 53 | 6 | AR428978 | Sequence |
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| CQ541527 | Sequence | 60 | 6 | CQ541527 | Sequence |
| AX786025 | Sequence | 20 | 6 | AX786025 | Sequence |
| BD262353 | Information | 26 | 6 | BD262353 | Information |
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| AX037776 | Sequence | 26 | 6 | AX037776 | Sequence |
| AX037777 | Sequence | 26 | 6 | AX037777 | Sequence |
| AR161742 | Sequence | 28 | 6 | AR161742 | Sequence |
| BD012790 | An animal | 28 | 6 | BD012790 | An animal |
| AR112406 | Sequence | 29 | 6 | AR112406 | Sequence |
| AX440319 | Sequence | 31 | 6 | AX440319 | Sequence |
| AR026654 | Sequence | 34 | 6 | AR026654 | Sequence |
| AX175152 | Sequence | 36 | 6 | AX175152 | Sequence |
| AR129977 | Sequence | 42 | 6 | AR129977 | Sequence |
| AR156107 | Sequence | 42 | 6 | AR156107 | Sequence |
| AR166385 | Sequence | 42 | 6 | AR166385 | Sequence |
| AR205069 | Sequence | 42 | 6 | AR205069 | Sequence |
| AR222034 | Sequence | 42 | 6 | AR222034 | Sequence |
| AR381164 | Sequence | 42 | 6 | AR381164 | Sequence |
| AR452252 | Sequence | 42 | 6 | AR452252 | Sequence |
| AR239886 | Sequence | 49 | 6 | AR239886 | Sequence |
| AX279688 | Sequence | 49 | 6 | AX279688 | Sequence |
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| AX116437 | Sequence | 51 | 6 | AX116437 | Sequence |
| AX202415 | Sequence | 51 | 6 | AX202415 | Sequence |
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| CQ539935 | Sequence | 60 | 6 | CQ539935 | Sequence |
| CQ562233 | Sequence | 60 | 6 | CQ562233 | Sequence |
| AX687679 | Sequence | 17 | 6 | AX687679 | Sequence |
| AX687680 | Sequence | 17 | 6 | AX687680 | Sequence |
| AX687681 | Sequence | 17 | 6 | AX687681 | Sequence |
| AX687682 | Sequence | 17 | 6 | AX687682 | Sequence |
| AX687683 | Sequence | 17 | 6 | AX687683 | Sequence |
| AX689176 | Sequence | 25 | 6 | AX689176 | Sequence |
| AX689177 | Sequence | 25 | 6 | AX689177 | Sequence |
| AX689178 | Sequence | 25 | 6 | AX689178 | Sequence |
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| AX689180 | Sequence | 25 | 6 | AX689180 | Sequence |
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| AX689186 | Sequence | 25 | 6 | AX689186 | Sequence |
| AX689187 | Sequence | 25 | 6 | AX689187 | Sequence |
| AX689188 | Sequence | 25 | 6 | AX689188 | Sequence |
| BD002867 | Gene comp | 31 | 6 | BD002867 | Gene comp |
| AX700433 | Sequence | 32 | 6 | AX700433 | Sequence |
| BD174127 | Lectin li | 33 | 6 | BD174127 | Lectin li |
| BD174334 | Lectin su | 33 | 6 | BD174334 | Lectin su |
| AX518053 | Sequence | 41 | 6 | AX518053 | Sequence |
| AX384367 | Sequence | 57 | 6 | AX384367 | Sequence |
| CQ544405 | Sequence | 60 | 6 | CQ544405 | Sequence |
| AR174416 | Sequence | 20 | 6 | AR174416 | Sequence |
| I71425 | Sequence 15 | 20 | 6 | I71425 | Sequence 15 |
| AR442472 | Sequence | 20 | 6 | AR442472 | Sequence |
| BD016108 | Oligonuc1 | 20 | 6 | BD016108 | Oligonuc1 |
| BD016227 | Oligonuc1 | 20 | 6 | BD016227 | Oligonuc1 |
| BD017379 | Oligonuc1 | 20 | 6 | BD017379 | Oligonuc1 |
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| AR090833 | Sequence | 23 | 6 | AR090833 | Sequence |
| BD249651 | Pi-ta gen | 23 | 6 | BD249651 | Pi-ta gen |
| AR197868 | Sequence | 23 | 6 | AR197868 | Sequence |
| AR254324 | Sequence | 23 | 6 | AR254324 | Sequence |
| AX260022 | Sequence | 23 | 6 | AX260022 | Sequence |
| AX038353 | Sequence | 23 | 6 | AX038353 | Sequence |
| AR080391 | Sequence | 24 | 6 | AR080391 | Sequence |
| AX038354 | Sequence | 24 | 6 | AX038354 | Sequence |
| AX180432 | Sequence | 24 | 6 | AX180432 | Sequence |
| AX445439 | Sequence | 24 | 6 | AX445439 | Sequence |

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c 94 11.2 58.9 26 6 AX038356
c 95 11.2 58.9 27 6 AX038557
c 96 11.2 58.9 27 6 AX597441
c 97 11.2 58.9 29 6 AX149585
c 98 11.2 58.9 29 6 I07501
c 99 11.2 58.9 29 6 AR368852
c 100 11.2 58.9 32 6 AX951643
c 101 11.2 58.9 34 6 BD141418
c 102 11.2 58.9 39 6 AX711338
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c 104 11.2 58.9 39 10 MMU232739
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c 106 11.2 58.9 41 6 AX517530
c 107 11.2 58.9 41 6 AX518427
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c 109 11.2 58.9 43 6 AX180641
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c 139 11 57.9 24 6 AR230172
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c 148 11 57.9 29 6 AR307624
c 149 11 57.9 29 6 AR371864
c 150 11 57.9 30 6 BD165838

ALIGNMENTS

RESULT 1
AR430777 AR430777 Sequence 80 from patent US 6649409. linear PAT 18-DEC-2003
LOCUS AR430777 24 bp DNA
DEFINITION AR430777
ACCESSION AR430777
VERSION AR430777.1 GI:40191706
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
Unclassified.

REFERENCE 1 (bases 1 to 24)
AUTHORS Fomesgaard, A.
TITLE Method for producing a nucleotide sequence construct with optimized codons for an HIV genetic vaccine based on a primary, early HIV isolate and synthetic envelope BX08 constructs
JOURNAL Patent: US 6649409-A 80 18-NOV-2003;
FEATURES
source Location/Qualifiers
1..24
/organism="unknown"
/mol_type="genomic DNA"
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Query Match 66.3%; Score 12.6; DB 6; Length 24;
Best Local Similarity 78.9%; Pred. No. 3.4e+05;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 1 CGGAATCCCCCGCGTGTG 19
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Db 2 CGGAATTCGCCCGCGTGTG 20
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RESULT 2
AX456376/c
LOCUS AX456376 54 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 234 from Patent WO0216944.
ACCESSION AX456376
VERSION AX456376.1 GI:21715280
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1
AUTHORS Wood, K.V., Wood, M.G., Zhuang, Y. and Paguio, A.
TITLE Synthetic nucleic acid molecule compositions and methods of preparation
JOURNAL Patent: WO 0216944-A 234 28-FEB-2002;
FEATURES
source Location/Qualifiers
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/organism="synthetic construct"
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/db_xref="taxon:32630"
/note="A primer"
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Qy 1 CGGAATCCCCCGCGTGTG 19
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Db 19 CGGAATGCCCAAGCTTTTG 1
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RESULT 3
AR300561/c
LOCUS AR300561 60 bp DNA linear PAT 12-JUN-2003
DEFINITION Sequence 12 from patent US 6537792.
ACCESSION AR300561
VERSION AR300561.1 GI:31688064
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
Unclassified.
REFERENCE 1 (bases 1 to 60)
AUTHORS Allen, M.J., Fang, T.-Y., Li, Y., Liu, H.-L., Chen, H.-M., Coutinho, P., Honzalko, R. and Ford, C.
TITLE Protein engineering of glucosylase to increase pH optimum, substrate specificity and thermostability
JOURNAL Patent: US 6537792-A 12 25-MAR-2003;
FEATURES
source Location/Qualifiers
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Query Match 64.2%; Score 12.2; DB 6; Length 31;
Best Local Similarity 73.7%; Pred. No. 5.2e+05;
Matches 14; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGTG 19
Db 19 CGGRCCTGCCCTGCATGGT 1

RESULT 8
ARI35209/c
LOCUS ARI35209 40 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 35 from patent US 6194559.
ACCESSION ARI35209
VERSION ARI35209.1 GI:14124114
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE Unclassified.
1 (bases 1 to 40)
AUTHORS Kim,S.Young.
TITLE Abscisic acid responsive element-binding transcription factors
JOURNAL Patent: US 6194559-A 35 27-FEB-2001;
FEATURES Location/Qualifiers
source 1..40
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/mol_type="unassigned DNA"

ORIGIN
Query Match 64.2%; Score 12.2; DB 6; Length 40;
Best Local Similarity 82.4%; Pred. No. 5e+05;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 GAATGCCCGCGTGTG 19
Db 40 GAATTCGCTCGTGTG 24

RESULT 9
ARI46705/c
LOCUS ARI46705 40 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 35 from patent US 6218527.
ACCESSION ARI46705
VERSION ARI46705.1 GI:15109894
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE Unclassified.
1 (bases 1 to 40)
AUTHORS Kim,S.Young.
TITLE Nucleic acid molecule encoding abscisic acid responsive element-binding factor 3
JOURNAL Patent: US 6218527-A 35 17-APR-2001;
FEATURES Location/Qualifiers
source 1..40
/organism="unknown"
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ORIGIN
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Best Local Similarity 82.4%; Pred. No. 5e+05;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 GAATGCCCGCGTGTG 19
Db 40 GAATTCGCTCGTGTG 24

RESULT 10
ARI52276/c
LOCUS ARI52276 40 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 35 from patent US 6232461.
ACCESSION ARI52276

VERSION ARI52276.1 GI:15118326
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE Unclassified.
1 (bases 1 to 40)
AUTHORS Kim,S.Young.
TITLE Nucleic acid molecule encoding abscisic acid responsive element-binding factor 4
JOURNAL Patent: US 6232461-A 35 15-MAY-2001;
FEATURES Location/Qualifiers
source 1..40
/organism="unknown"
/mol_type="unassigned DNA"

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Query Match 64.2%; Score 12.2; DB 6; Length 40;
Best Local Similarity 82.4%; Pred. No. 5e+05;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 GAATGCCCGCGTGTG 19
Db 40 GAATTCGCTCGTGTG 24

RESULT 11
ARI57814/c
LOCUS ARI57814 40 bp DNA linear PAT 17-OCT-2001
DEFINITION Sequence 35 from patent US 6245905.
ACCESSION ARI57814
VERSION ARI57814.1 GI:16218827
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE Unclassified.
1 (bases 1 to 40)
AUTHORS Kim,S.Young.
TITLE Nucleic acid molecule encoding abscisic acid responsive element-binding factor 2
JOURNAL Patent: US 6245905-A 35 12-JUN-2001;
FEATURES Location/Qualifiers
source 1..40
/organism="unknown"
/mol_type="unassigned DNA"

ORIGIN
Query Match 64.2%; Score 12.2; DB 6; Length 40;
Best Local Similarity 82.4%; Pred. No. 5e+05;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 GAATGCCCGCGTGTG 19
Db 40 GAATTCGCTCGTGTG 24

RESULT 12
CQ005341
LOCUS CQ005341 50 bp DNA linear PAT 16-JAN-2004
DEFINITION Sequence 3981 from Patent WO0147944.
ACCESSION CQ005341
VERSION CQ005341.1 GI:41011973
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
1 Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0147944-A 3981 05-JUL-2001;
FEATURES Curagen Corporation (US)
Location/Qualifiers

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Accession number c943328330"

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Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GGAATGCCCGCGTGT 18
Db 14 GCAATTCGCCGCGTGT 30

RESULT 13
AR523798/c
LOCUS AR523798 55 bp DNA linear PAT 22-SEP-2004
DEFINITION Sequence 28758 from patent US 6703491.
ACCESSION AR523798
VERSION AR523798.1 GI:52459273
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 55)
Homburger,S.A., Ebens,A.J. Jr., Erickson,C.S., Francis-Lang,H.L.,
Margolis,J.S., Reddy,B.P., Ruddy,D.A. and Buchman,A.R.
TITLE Drosophila sequences
JOURNAL Patent: US 6703491-A 28758 09-MAR-2004;
FEATURES
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Location/Qualifiers
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ORIGIN

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Best Local Similarity 64.2%; Score 12.2; DB 6; Length 55;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 GAATGCCCGCGTGT 19
Db 29 GCATGCCCGCGTGT 13

RESULT 14
AX772391
LOCUS AX772391 49 bp DNA linear PAT 02-JUL-2003
DEFINITION Sequence 181 from Patent WO03042407.
ACCESSION AX772391
VERSION AX772391.1 GI:32438964
KEYWORDS
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster
AUTHORS Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;
TITLE Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
JOURNAL Ephynroidea; Drosophilidae; Drosophila.
REFERENCE
1
Dickson,B., Berger,J., Suzuki,T. and Knoblich,J.
METHOD for identifying therapeutic targets by use of genetic
screens in drosophila melanogaster
PATENT: WO 03042407-A 181 22-MAY-2003;
BOEHRINGER INGELHEIM INTERNATIONAL GMBH; CD Patents (DE)
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Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GGAATGCCCGCG 13
Db 10 GGAATGCCCGCG 21

RESULT 15
CQ840727/c
LOCUS CQ840727 20 bp DNA linear PAT 29-JUL-2004
DEFINITION Sequence 2 from Patent WO2004058999.
ACCESSION CQ840727
VERSION CQ840727.1 GI:50838338
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE
1
Zimmermann,G. and Alexander,H.
METHOD and means for determining specific conditions or changes in
the uterine epithelium and in the epithelium of other organs
PATENT: WO 2004058999-A 2 15-JUL-2004;
JOURNAL Universitaet Leipzig (DE)
FEATURES
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Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 5 ATGCCCGCGTGTG 19
Db 17 ATGACCGCGTGTG 3

RESULT 16
AX155207
LOCUS AX155207 24 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 3 from Patent WO0140273.
ACCESSION AX155207
VERSION AX155207.1 GI:14536688
KEYWORDS
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
JOURNAL Cypriniformes; Cyprinidae; Danio.
REFERENCE
1
Uckun,F.M.
TITLE Transgenic zebra fish embryo model for hematopoiesis and
lymphoproliferative disorders
PATENT: WO 0140273-A 3 07-JUN-2001;
Parker Hughes Institute (US)
FEATURES
source
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Location/Qualifiers
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/db_xref="taxon:7955"

ORIGIN

Query Match
Best Local Similarity 62.1%; Score 11.8; DB 6; Length 24;
Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 5 ATGCCCGCGTGTG 19
Db 17 ATGACCGCGTGTG 19
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Db 5 ATGCCCTCGTGTG 19

RESULT 17
LOCUS A97649 48 bp DNA linear PAT 26-JAN-2000
DEFINITION Sequence 16 from Patent WO9915549.
ACCESSION A97649
VERSION A97649.1 GI:6780936
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 48)
AUTHORS Humphreys,D.P.
TITLE
JOURNAL
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Location/Qualifiers
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Best Local Similarity 86.7%; Pred. No. 7.6e+05;
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Qy 5 ATGCCCGCGTGTG 19
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Db 21 ATGCCCGCGTGTG 35

RESULT 18
LOCUS AR428977 48 bp DNA linear PAT 18-DEC-2003
DEFINITION Sequence 16 from patent US 6642356.
ACCESSION AR428977
VERSION AR428977.1 GI:40189018
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 48)
AUTHORS Humphreys,D.P.
TITLE Peptides which function as hinge regions in protein
JOURNAL Patent: US 6642356-A 16 04-NOV-2003;
FEATURES
source
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Location/Qualifiers
/organism="unknown"
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ORIGIN
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Best Local Similarity 86.7%; Pred. No. 7.6e+05;
Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 5 ATGCCCGCGTGTG 19
|||||
Db 21 ATGCCCGCGTGTG 35

RESULT 19
LOCUS BD081713 48 bp DNA linear PAT 27-AUG-2002
DEFINITION Peptide.
ACCESSION BD081713
VERSION BD081713.1 GI:22627316
KEYWORDS JP 2001517423-A/5.
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 48)
AUTHORS Humphreys,D.P.
TITLE Peptides which function as hinge regions in protein
JOURNAL Patent: US 6642356-A 16 04-NOV-2003;
FEATURES
source
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Location/Qualifiers
/organism="unassigned DNA"
/mol_type="genomic DNA"
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ORIGIN
Query Match 62.1%; Score 11.8; DB 6; Length 48;
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Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 5 ATGCCCGCGTGTG 19
|||||
Db 21 ATGCCCGCGTGTG 35

RESULT 20
LOCUS AX165870 50 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 1065 from Patent WO0138586.
ACCESSION AX165870
VERSION AX165870.1 GI:14546699
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
REFERENCE 1
AUTHORS Shimkets R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and
JOURNAL Patent: WO 0138586-A 1065 31-MAY-2001;
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misc_feature
variation

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Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GGAATGCCCGCGTGTG 16
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Db 23 GGAATGCCCGCGTGTG 37

RESULT 21
LOCUS A97650 53 bp DNA linear PAT 26-JAN-2000
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 48)
AUTHORS Humphreys,D.P.
TITLE
JOURNAL
FEATURES
source
1..48
Location/Qualifiers
/organism="Artificial Sequence".
/mol_type="synthetic construct"
/db_xref="taxon:32630"

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Best Local Similarity 86.7%; Pred. No. 7.6e+05;
Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 5 ATGCCCGCGTGTG 19
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Db 21 ATGCCCGCGTGTG 35
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DEFINITION Sequence 17 from Patent WO9915549.
ACCESSION A97650
VERSION A97650.1 GI:6780937
SOURCE .
KEYWORDS unidentified
ORGANISM unidentified
REFERENCE 1 (bases 1 to 53)
AUTHORS Humphreys,D.P.
TITLE PEPTIDES
JOURNAL Patent: WO 9915549-A 17 01-APR-1999;
CELLTECH THERAPEUTICS LTD (GB); HUMPHREYS DAVID PAUL (GB)
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Qy 5 ATGCCCGCGGTGG 19
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Db 20 ATGCCCGCGGTGG 34

RESULT 22
AR428978 LOCUS AR428978 53 bp DNA linear PAT 18-DEC-2003
DEFINITION Sequence 17 from patent US 6642356.
ACCESSION AR428978
VERSION AR428978.1 GI:40189019
KEYWORDS Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 53)
AUTHORS Humphreys,D.P.
TITLE Peptides which function as hinge regions in protein
JOURNAL Patent: US 6642356-A 17 04-NOV-2003;
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Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 5 ATGCCCGCGGTGG 19
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Db 20 ATGCCCGCGGTGG 34

RESULT 23
BD081714 LOCUS BD081714 53 bp DNA linear PAT 27-AUG-2002
DEFINITION Peptide.
ACCESSION BD081714
VERSION BD081714.1 GI:32627317
KEYWORDS JP 2001517423-A/6.
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 53)
AUTHORS Humphreys,D.P.
TITLE Peptide
JOURNAL Patent: JP 2001517423-A 6 09-OCT-2001;
CELLTECH THERAPEUTICS LTD
COMMENT OS Artificial Sequence

PN JP 2001517423-A/6
PD 09-OCT-2001
PF 21-SEP-1998 JP 2000512854
PR 19-SEP-1997 GB 9720054.7
PI DAVID PAUL HUMPHREYS
PC C12N15/09,C07K7/08,C07K16/00,C12N15/00
CC double stranded except for the four 5' residues on each strand
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Best Local Similarity 86.7%; Pred. No. 7.5e+05;
Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 20 ATGCCCGCGGTGG 34

RESULT 24
CQ541527 LOCUS CQ541527 60 bp DNA linear PAT 30-JAN-2004
DEFINITION Sequence 11162 from Patent WO0210449.
ACCESSION CQ541527
VERSION CQ541527.1 GI:41507791
KEYWORDS Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Shoshan,A., Wasserman,A., Mintz,E., Mintz,I. and Faigler,S.
TITLE Oligonucleotide library for detecting rna transcripts and splice
variants that populate a transcriptome
JOURNAL Patent: WO 0210449-A 1162 07-FEB-2002;
CompuGen Inc. (US)
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Db 24 AATGCTCTGCGGTGT 38

RESULT 25
AX786025 LOCUS AX786025 20 bp DNA linear PAT 17-JUL-2003
DEFINITION Sequence 21 from Patent WO03050272.
ACCESSION AX786025
VERSION AX786025.1 GI:32953645
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Bandelier,M.A., Denys,P., Denormandie,P., Sapena,R.,
Lepailleur-Enouf,D. and Youssefian,T.
TITLE Bone development model
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JOURNAL Patent: WO 03050272-A 21 19-JUN-2003;
Sympathos (FR)
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Db 1 CGGAATGCCCTGTGTT 18
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BD262353
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DEFINITION Informationcarrying and -processing polymers.
ACCESSION BD262353
VERSION BD262353.1 GI:33072121
KEYWORDS JP 2002541539-A/127.
SOURCE synthetic construct
ORGANISM
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REFERENCE
  1 (bases 1 to 26)
  Raue,H., Feldkamp,U., Banzhaf,W. and Howard,J.C.
  Informationcarrying and -processing polymers
  Patent: JP 2002541539-A 127 03-DEC-2002;
  HILMAR RAUHE
TITLE Informationcarrying and -processing polymers
JOURNAL
  Patent: JP 2002541539-A 127 03-DEC-2002;
  HILMAR RAUHE
COMMENT
  OS Artificial Sequence
  PN JP 2002541539-A/127
  PD 03-DEC-2002
  PF 31-MAR-2000 JP 2000609427
  PR 31-MAR-1999 DE 199 14 808.2
  PI HILMAR RAUHE,UDO FELDKAMP,WOLFGANG BANZHAF,JONATHAN C HOWARD
  PC G06N3/12,C12M1/00,C12N15/09,C12N15/09,C12Q1/68,C12Q1/68,G06N1/
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Db 23 GTAATACCACGGGTGTTG 6
RESULT 28
AX037776
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DEFINITION Sequence 127 from Patent WO0059917.
ACCESSION AX037776
VERSION AX037776.1 GI:11227158
KEYWORDS
SOURCE synthetic construct
ORGANISM
  other sequences; artificial sequences.
REFERENCE
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  Howard,J.C., Feldkamp,U., Raue,H. and Banzhaf,W.
  Information-carrying and information-processing polymers
  Patent: WO 0059917-A 127 12-OCT-2000;
  HOWARD JONATHAN C (DE) ; FELDKAMP UDO (DE) ; RAUHE HILMAR (DE) ;
  BANZHAF WOLFGANG (DE)
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Db 8 GTAATACCACGGGTGTTG 25
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DEFINITION Informationcarrying and -processing polymers.
ACCESSION BD262354
VERSION BD262354.1 GI:33072122
KEYWORDS JP 2002541539-A/128.
SOURCE synthetic construct
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REFERENCE
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  Raue,H., Feldkamp,U., Banzhaf,W. and Howard,J.C.
  Informationcarrying and -processing polymers
  Patent: JP 2002541539-A 128 03-DEC-2002;
  HILMAR RAUHE
COMMENT
  OS Artificial Sequence
  PN JP 2002541539-A/128
  PD 03-DEC-2002
  PF 31-MAR-2000 JP 2000609427
  PR 31-MAR-1999 DE 199 14 808.2
  PI HILMAR RAUHE,UDO FELDKAMP,WOLFGANG BANZHAF,JONATHAN C HOWARD
  PC G06N3/12,C12M1/00,C12N15/09,C12N15/09,C12Q1/68,C12Q1/68,G06N1/
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  PC C07H21/00,C12N15/00,C12N15/00
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  data.
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Qy 2 GGAATGCCCGCGGTGTTG 19
Db 23 GTAATACCACGGGTGTTG 6
RESULT 28
AX037776
LOCUS 26 bp DNA linear PAT 16-NOV-2000
DEFINITION Sequence 127 from Patent WO0059917.
ACCESSION AX037776
VERSION AX037776.1 GI:11227158
KEYWORDS
SOURCE synthetic construct
ORGANISM
  other sequences; artificial sequences.
REFERENCE
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  Howard,J.C., Feldkamp,U., Raue,H. and Banzhaf,W.
  Information-carrying and information-processing polymers
  Patent: WO 0059917-A 127 12-OCT-2000;
  HOWARD JONATHAN C (DE) ; FELDKAMP UDO (DE) ; RAUHE HILMAR (DE) ;
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Db 8 GTAATACCACGGGTGTTG 25
RESULT 29
AX037777/c
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LOCUS AX037777 26 bp DNA linear PAT 16-NOV-2000
DEFINITION Sequence 128 from Patent WO0059917.
ACCESSION AX037777
VERSION AX037777.1 GI:11227159
KEYWORDS

SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1
AUTHORS Howard, J.C., Feldkamp, U., Rauhe, H. and Banzhaf, W.
TITLE Information-carrying and information-processing polymers
JOURNAL Patent: WO 0059917-A 128 12-OCT-2000; RAUHE HILMAR (DE);
HOWARD JONATHAN C (DE); FELDKAMP UDO (DE);
BANZHAF WOLFGANG (DE)

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Matches 14; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 2 GGAATGCCCGCGGTG 19
Db 23 GTATACACCGGTG 6

RESULT 30
AR161742/c

LOCUS AR161742 28 bp DNA linear PAT 17-OCT-2001
DEFINITION Sequence 52 from patent US 6258529.
ACCESSION AR161742
VERSION AR161742.1 GI:16228657
KEYWORDS

SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 28)
AUTHORS Berdoz, J. and Kraehenbuhl, J.-P.
TITLE PCR amplification of rearranged genomic variable regions of immunoglobulin genes
JOURNAL Patent: US 6258529-A 52 10-JUL-2001;

FEATURES
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Matches 14; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 2 GGAATGCCCGCGGTG 19
Db 24 GCAATGCCCTGTGTGTCG 7

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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Title: US-10-729-421-52

Perfect score: 19
Sequence: 1 cqaatgccccgcgtattg 19

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Maximum Match 100%
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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| 3 | 17 | 89.5 | 17 | 6 | ACN01440 | Acn01440 WNV Inozv |
| 4 | 17 | 89.5 | 17 | 6 | ACN01439 | Acn01439 WNV Inozv |
| C 5 | 17 | 89.5 | 17 | 6 | ACN15269 | Acn15269 WNV minus |
| 6 | 17 | 89.5 | 17 | 6 | ACN01438 | Acn01438 WNV Inozv |
| C 7 | 17 | 89.5 | 17 | 6 | ACN15270 | Acn15270 WNV minus |
| C 8 | 16 | 84.2 | 17 | 6 | ACN13601 | Acn13601 WNV minus |
| 9 | 16 | 84.2 | 17 | 6 | ACN03481 | Acn03481 WNV Zinzv |
| C 10 | 15 | 78.9 | 17 | 6 | ACN03600 | Acn03600 WNV minus |
| 11 | 15 | 78.9 | 17 | 6 | ACN03480 | Acn03480 WNV Zinzv |
| C 12 | 15 | 78.9 | 23 | 12 | ADN36699 | Adn36699 West Nile |
| C 13 | 15 | 78.9 | 50 | 12 | ADN36711 | Adn36711 West Nile |
| C 14 | 14.2 | 74.7 | 19 | 12 | ADQ30683 | Adq30683 West Nile |
| 15 | 14.2 | 74.7 | 19 | 12 | ADQ30679 | Adq30679 West Nile |
| 16 | 14 | 73.7 | 17 | 6 | ACN03482 | Acn03482 WNV Zinzv |
| C 17 | 14 | 73.7 | 17 | 6 | ACN12235 | Acn12235 WNV minus |
| C 18 | 14 | 73.7 | 23 | 12 | ADN36700 | Adn36700 West Nile |
| C 19 | 14 | 73.7 | 50 | 12 | ADN36712 | Adn36712 West Nile |
| C 20 | 13 | 68.4 | 17 | 6 | ACN04726 | Acn04726 WNV DNZv |

| | | | | | |
|-------|------|------|----|----|----------|
| 94 | 11.6 | 61.1 | 42 | 6 | AAK98822 |
| c 95 | 11.6 | 61.1 | 42 | 10 | ACC44780 |
| 96 | 11.6 | 61.1 | 43 | 10 | ACC44779 |
| 97 | 11.6 | 61.1 | 43 | 12 | ADF82849 |
| 98 | 11.6 | 61.1 | 43 | 12 | ADF82845 |
| 99 | 11.6 | 61.1 | 43 | 12 | ADF82851 |
| 100 | 11.6 | 61.1 | 43 | 13 | ADR73321 |
| 101 | 11.6 | 61.1 | 46 | 10 | ADF42963 |
| 102 | 11.6 | 61.1 | 46 | 10 | ADL18079 |
| 103 | 11.6 | 61.1 | 49 | 5 | ABA10731 |
| c 104 | 11.6 | 61.1 | 50 | 4 | AHH89661 |
| c 105 | 11.6 | 61.1 | 51 | 4 | ADL11577 |
| 106 | 11.6 | 61.1 | 51 | 4 | AHH89661 |
| 107 | 11.6 | 61.1 | 60 | 6 | ABN59120 |
| 108 | 11.6 | 61.1 | 60 | 6 | ABN36822 |
| 109 | 11.4 | 60.0 | 17 | 8 | ADA99424 |
| 110 | 11.4 | 60.0 | 17 | 8 | ADA99423 |
| 111 | 11.4 | 60.0 | 17 | 8 | ADA99426 |
| 112 | 11.4 | 60.0 | 17 | 8 | ADA99425 |
| 113 | 11.4 | 60.0 | 17 | 8 | ADA99422 |
| 114 | 11.4 | 60.0 | 18 | 6 | AD38567 |
| 115 | 11.4 | 60.0 | 20 | 13 | ADR67378 |
| 116 | 11.4 | 60.0 | 25 | 8 | ADB00923 |
| 117 | 11.4 | 60.0 | 25 | 8 | ADB00929 |
| 118 | 11.4 | 60.0 | 25 | 8 | ADB00925 |
| 119 | 11.4 | 60.0 | 25 | 8 | ADB00932 |
| 120 | 11.4 | 60.0 | 25 | 8 | ADB00926 |
| 121 | 11.4 | 60.0 | 25 | 8 | ADB00927 |
| 122 | 11.4 | 60.0 | 25 | 8 | ADB00928 |
| 123 | 11.4 | 60.0 | 25 | 8 | ADB00931 |
| 124 | 11.4 | 60.0 | 25 | 8 | ADB00930 |
| 125 | 11.4 | 60.0 | 25 | 8 | ADB00922 |
| 126 | 11.4 | 60.0 | 25 | 8 | ADB00924 |
| 127 | 11.4 | 60.0 | 25 | 8 | ADB00934 |
| 128 | 11.4 | 60.0 | 25 | 8 | ADB00933 |
| c 129 | 11.4 | 60.0 | 25 | 9 | ACH53197 |
| c 130 | 11.4 | 60.0 | 31 | 2 | ACH53197 |
| c 131 | 11.4 | 60.0 | 31 | 3 | AA79163 |
| c 132 | 11.4 | 60.0 | 32 | 10 | ABX94700 |
| c 133 | 11.4 | 60.0 | 33 | 6 | ABS67932 |
| c 134 | 11.4 | 60.0 | 33 | 6 | ABS68670 |
| c 135 | 11.4 | 60.0 | 33 | 12 | ADL95719 |
| c 136 | 11.4 | 60.0 | 57 | 6 | ABK11619 |
| c 137 | 11.4 | 60.0 | 57 | 6 | ACN24584 |
| c 138 | 11.4 | 60.0 | 60 | 6 | ABN41292 |
| c 139 | 11.2 | 58.9 | 18 | 10 | ADF18572 |
| 140 | 11.2 | 58.9 | 18 | 10 | ADF18575 |
| 141 | 11.2 | 58.9 | 20 | 2 | AAQ25130 |
| 142 | 11.2 | 58.9 | 20 | 2 | AAQ84274 |
| c 143 | 11.2 | 58.9 | 20 | 4 | AAF23206 |
| c 144 | 11.2 | 58.9 | 20 | 6 | ABA02264 |
| c 145 | 11.2 | 58.9 | 20 | 12 | ADK96038 |
| c 146 | 11.2 | 58.9 | 21 | 4 | AAF96169 |
| c 147 | 11.2 | 58.9 | 21 | 10 | ADD20604 |
| 148 | 11.2 | 58.9 | 22 | 10 | AD63716 |
| 149 | 11.2 | 58.9 | 22 | 10 | ADL01698 |
| c 150 | 11.2 | 58.9 | 23 | 2 | AAT43862 |

ALIGNMENTS

RESULT 1
ADQ30682
ID ADQ30682 standard; DNA; 19 BP.
XX
XX ADQ30682;
XX
DT 23-SEP-2004 (first entry)
XX
XX West Nile Virus oligonucleotide probe A.
DE
XX ss; probe; West Nile Virus; diagnosis.

XX West Nile virus.
OS WO2004055159-A2.
PN 01-JUL-2004.
XX 05-DEC-2003; 2003WO-US038750.
PR 12-DEC-2002; 2002US-0432850P.
PR 20-JUN-2003; 2003US-0480431P.
(CHIR) CHIRON CORP.
PI Shyamala V;
XX WPI; 2004-488058/46.
XX New isolated oligonucleotides for accurately diagnosing West Nile virus infection or for capturing, detecting and quantitating West Nile virus in blood samples.
XX Claim 1; SEQ ID NO 52; 56pp; English.
XX The invention relates to an isolated oligonucleotide not more than 60 nucleotides in length comprising a nucleotide sequence (S1) of at least 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g. 20, 21 or 23 bp) given in the specification derived from the West Nile virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence identity to the nucleotide sequence of (S1), or complements of (S1) and (S2). The oligonucleotide further comprises a detectable label at the 5'-end and/or the 3'-end. The detectable label is a fluorescent label selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine (TAMRA), and 2',4',5',7',-tetrachloro-4,7-dichlorofluorescein (TET). The composition and methods are useful for accurately diagnosing West Nile virus infection or for capturing, detecting and quantitating West Nile virus in biological samples, particularly blood samples. This sequence corresponds to an oligonucleotide probe of the invention.
XX Sequence 19 BP; 2 A; 6 C; 7 G; 4 T; 0 U; 0 Other;
XX Query Match 100.0%; Score 19; DB 12; Length 19;
XX Best Local Similarity 100.0%; Pred. No. 5.5;
XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CGGAATGCCCGCGTGTG 19
Db 1 CGGAATGCCCGCGTGTG 19
RESULT 2
ACN15268/C
ID ACN15268 standard; RNA; 17 BP.
XX ACN15268;
XX 22-APR-2004 (first entry)
XX WNV minus strand Ambenzyme substrate SEQ ID NO 15271.
DE WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic; virucide; neuroprotective; antibacterial; replication; pancreatitis; encephalitis; myocarditis; meningitis; infection; hepatitis; liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNazyme; Ambenzyme; Zinzyne; ss.
XX West Nile Virus.
OS
XX WO200268637-A2.
XX
XX 06-SEP-2002.
XX
XX 19-OCT-2001; 2001WO-US048350.

CC are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
CC in the specification. The present sequence is that of a nucleic acid
CC molecule of the invention

XX Sequence 17 BP; 2 A; 6 C; 6 G; 0 T; 3 U; 0 Other;

Query Match 89.5%; Score 17; DB 6; Length 17;

Best Local Similarity 82.4%; Pred. No. 59;

Matches 14; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGT 17

Db 1 CGGAATGCCCGCGTGT 17

RESULT 7

ACN15270/c

ID ACN15270 standard; RNA; 17 BP.

XX AC

XX ACN15270;

XX 22-APR-2004 (first entry)

XX DE

XX WNV minus strand Amberzyme substrate SEQ ID NO 15273.

XX OS

XX WNV; West Nile Virus; antiinflammatory; cytosolic; hepatotropic;

XX KW virucide; neuroprotective; antibacterial; replication; pancreatitis;

XX KW encephalitis; myocarditis; meningitis; infection; hepatitis;

XX KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;

XX KW Amberzyme; Zinzyme; ss.

XX XX

XX West Nile Virus.

XX OS

XX WO200268637-A2.

XX PN

XX 06-SEP-2002.

XX PD

XX 19-OCT-2001; 2001WO-US048350.

XX PF

XX 20-OCT-2000; 2000US-0242411P.

XX PR

XX (RIBO-) RIBOZYME PHARM INC.

XX PA (BLAT/) BLATT L.

XX PA (MCSW/) MCSWIGGEN J A.

XX PI Blatt L, Mcswiggen JA;

XX XX

XX WPI; 2002-706994/76.

XX DR

XX New nucleic acid molecule that modulates replication of West Nile Virus

XX PT (WNV), useful for treating a condition related to WNV infection e.g.

XX PT pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.

XX XX

XX Claim 23; SEQ ID NO 15273; 495pp; English.

XX PS

XX The invention relates to nucleic acid molecules that modulate replication

XX CC of the West Nile Virus (WNV). The nucleic acid molecules are useful for

XX CC treating a condition related to WNV infection e.g. pancreatitis,

XX CC encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,

XX CC liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid

XX CC molecule is selected from the group of ribozymes consisting of

XX CC Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyme. The

XX CC nucleic acid molecules further comprise at least five ribose residues, at

XX CC least ten 2'-O-methyl modifications, phosphorothioate linkages on at

XX CC least three of the 5' terminal nucleotides and a 3' end modification of a

XX CC 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080

XX CC are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given

XX CC in the specification. The present sequence is that of a nucleic acid

XX CC molecule of the invention

XX SQ

Sequence 17 BP; 3 A; 6 C; 6 G; 0 T; 2 U; 0 Other;

Query Match 89.5%; Score 17; DB 6; Length 17;

Best Local Similarity 100.0%; Pred. No. 59;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGT 17

Db 17 CGGAATGCCCGCGTGT 1

RESULT 8

ACN13601/c

ID ACN13601 standard; RNA; 17 BP.

XX AC

XX ACN13601;

XX XX

XX 22-APR-2004 (first entry)

XX DT

XX WNV minus strand Zinzyme substrate SEQ ID NO 13604.

XX DE

XX WNV; West Nile Virus; antiinflammatory; cytosolic; hepatotropic;

XX KW virucide; neuroprotective; antibacterial; replication; pancreatitis;

XX KW encephalitis; myocarditis; meningitis; infection; hepatitis;

XX KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;

XX KW Amberzyme; Zinzyme; ss.

XX XX

XX West Nile Virus.

XX OS

XX WO200268637-A2.

XX PN

XX 06-SEP-2002.

XX PD

XX 19-OCT-2001; 2001WO-US048350.

XX PF

XX 20-OCT-2000; 2000US-0242411P.

XX PR

XX (RIBO-) RIBOZYME PHARM INC.

XX PA (BLAT/) BLATT L.

XX PA (MCSW/) MCSWIGGEN J A.

XX PI Blatt L, Mcswiggen JA;

XX XX

XX WPI; 2002-706994/76.

XX DR

XX New nucleic acid molecule that modulates replication of West Nile Virus

XX PT (WNV), useful for treating a condition related to WNV infection e.g.

XX PT pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.

XX XX

XX Claim 23; SEQ ID NO 13604; 495pp; English.

XX PS

XX The invention relates to nucleic acid molecules that modulate replication

XX CC of the West Nile Virus (WNV). The nucleic acid molecules are useful for

XX CC treating a condition related to WNV infection e.g. pancreatitis,

XX CC encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,

XX CC liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid

XX CC molecule is selected from the group of ribozymes consisting of

XX CC Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyme. The

XX CC nucleic acid molecules further comprise at least five ribose residues, at

XX CC least ten 2'-O-methyl modifications, phosphorothioate linkages on at

XX CC least three of the 5' terminal nucleotides and a 3' end modification of a

XX CC 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080

XX CC are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given

XX CC in the specification. The present sequence is that of a nucleic acid

XX CC molecule of the invention

XX SQ

Sequence 17 BP; 2 A; 7 C; 6 G; 0 T; 2 U; 0 Other;

Query Match 84.2%; Score 16; DB 6; Length 17;

Best Local Similarity 100.0%; Pred. No. 1.9e+02;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGT 16

Db 16 CGGAATGCCCGCGTGT 1

```
RESULT 9
ACN03481
ID ACN03481 standard; RNA; 17 BP.
AC ACN03481;
XX
DT 22-APR-2004 (first entry)
XX
DE WNV Zinzyne substrate SEQ ID NO 3484.
XX
WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
KW encephalitis; myocarditis; meningitis; infection; hepatitis;
KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;
KW Amberzyme; Zinzyne; ss.
XX
OS West Nile Virus.
XX
PN WO200268637-A2.
XX
PD 06-SEP-2002.
XX
PF 19-OCT-2001; 2001WO-US048350.
XX
PR 20-OCT-2000; 2000US-0242411P.
XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (BLAT/) BLATT L.
PA (MCSW/) MCSWIGGEN J A.
XX
PI Blatt L, Mcswiggen JA;
XX
WPI; 2002-706994/76.
XX
New nucleic acid molecule that modulates replication of West Nile Virus
(WNV), useful for treating a condition related to WNV infection e.g.
pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX
Claim 23; SEQ ID NO 3484; 495pp; English.
XX
The invention relates to nucleic acid molecules that modulate replication
of the West Nile Virus (WNV). The nucleic acid molecules are useful for
treating a condition related to WNV infection e.g. pancreatitis,
encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
molecule is selected from the group of ribozymes consisting of
Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyne. The
nucleic acid molecules further comprise at least five ribose residues, at
least ten 2'-O-methyl modifications, phosphorothioate linkages on at
least three of the 5' terminal nucleotides and a 3' end modification of a
3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
in the specification. The present sequence is that of a nucleic acid
molecule of the invention
XX
Sequence 17 BP; 2 A; 5 C; 5 G; 0 T; 5 U; 0 Other;
Query Match 84.2%; Score 16; DB 6; Length 17;
Best Local Similarity 75.0%; Pred. NO. 1.9e+02;
Matches 12; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
OY 4 AATGCCCGCGGTG 19
DB 1 AAUGCCCGCGUGUG 16
RESULT 10
ACN13600/c
ID ACN13600 standard; RNA; 17 BP.
XX
AC ACN13600;
XX
WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
KW encephalitis; myocarditis; meningitis; infection; hepatitis;
KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;
KW Amberzyme; Zinzyne; ss.
XX
OS West Nile Virus.
XX
PN WO200268637-A2.
XX
PD 06-SEP-2002.
XX
PF 19-OCT-2001; 2001WO-US048350.
XX
PR 20-OCT-2000; 2000US-0242411P.
XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (BLAT/) BLATT L.
PA (MCSW/) MCSWIGGEN J A.
XX
PI Blatt L, Mcswiggen JA;
XX
WPI; 2002-706994/76.
XX
New nucleic acid molecule that modulates replication of West Nile Virus
(WNV), useful for treating a condition related to WNV infection e.g.
pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX
Claim 23; SEQ ID NO 3484; 495pp; English.
XX
The invention relates to nucleic acid molecules that modulate replication
of the West Nile Virus (WNV). The nucleic acid molecules are useful for
treating a condition related to WNV infection e.g. pancreatitis,
encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
molecule is selected from the group of ribozymes consisting of
Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyne. The
nucleic acid molecules further comprise at least five ribose residues, at
least ten 2'-O-methyl modifications, phosphorothioate linkages on at
least three of the 5' terminal nucleotides and a 3' end modification of a
3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
in the specification. The present sequence is that of a nucleic acid
molecule of the invention
XX
Sequence 17 BP; 2 A; 5 C; 5 G; 0 T; 5 U; 0 Other;
Query Match 84.2%; Score 16; DB 6; Length 17;
Best Local Similarity 75.0%; Pred. NO. 1.9e+02;
Matches 12; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
OY 4 AATGCCCGCGGTG 19
DB 1 AAUGCCCGCGUGUG 16
RESULT 11
ACN03480
ID ACN03480 standard; RNA; 17 BP.
XX
AC ACN03480;
XX
22-APR-2004 (first entry)
XX
WNV Zinzyne substrate SEQ ID NO 3483.
XX
WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
KW encephalitis; myocarditis; meningitis; infection; hepatitis;
KW Amberzyme; Zinzyne; ss.
XX
OS West Nile Virus.
XX
PN WO200268637-A2.
XX
PD 06-SEP-2002.
XX
PF 19-OCT-2001; 2001WO-US048350.
XX
PR 20-OCT-2000; 2000US-0242411P.
XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (BLAT/) BLATT L.
PA (MCSW/) MCSWIGGEN J A.
XX
PI Blatt L, Mcswiggen JA;
XX
WPI; 2002-706994/76.
XX
New nucleic acid molecule that modulates replication of West Nile Virus
(WNV), useful for treating a condition related to WNV infection e.g.
pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX
Claim 23; SEQ ID NO 13603; 495pp; English.
XX
The invention relates to nucleic acid molecules that modulate replication
of the West Nile Virus (WNV). The nucleic acid molecules are useful for
treating a condition related to WNV infection e.g. pancreatitis,
encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
molecule is selected from the group of ribozymes consisting of
Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyne. The
nucleic acid molecules further comprise at least five ribose residues, at
least ten 2'-O-methyl modifications, phosphorothioate linkages on at
least three of the 5' terminal nucleotides and a 3' end modification of a
3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
in the specification. The present sequence is that of a nucleic acid
molecule of the invention
XX
Sequence 17 BP; 5 A; 5 C; 6 G; 0 T; 1 U; 0 Other;
Query Match 78.9%; Score 15; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. NO. 6.4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 ATGCCCGCGGTG 19
DB 17 ATGCCCGCGGTG 3
RESULT 11
ACN03480
ID ACN03480 standard; RNA; 17 BP.
XX
AC ACN03480;
XX
22-APR-2004 (first entry)
XX
WNV Zinzyne substrate SEQ ID NO 3483.
XX
WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
KW encephalitis; myocarditis; meningitis; infection; hepatitis;
```

KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNazyme;
 KW Amberyze; Zinzyme; ss.

XX
 XX West Nile Virus.

XX
 XX WO200268637-A2.

XX
 XX 06-SEP-2002.

XX
 XX 19-OCT-2001; 2001WO-US048350.

XX
 XX 20-OCT-2000; 2000US-0242411P.

XX
 XX (RIBO-) RIBOZYME PHARM INC.

XX
 XX (BLAT/) BLATT L.

XX
 XX (MCSW/) MCSWIGGEN J A.

XX
 XX Blatt L, Mcswiggen JA;

XX
 XX WPI; 2002-706994/76.

XX
 XX New nucleic acid molecule that modulates replication of West Nile Virus
 (WNV), useful for treating a condition related to WNV infection e.g.
 pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.

XX
 XX Claim 23; SEQ ID NO 3483; 495pp; English.

XX
 XX The invention relates to nucleic acid molecules that modulate replication
 of the West Nile Virus (WNV). The nucleic acid molecules are useful for
 treating a condition related to WNV infection e.g. pancreatitis,
 encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
 liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
 molecule is selected from the group of ribozymes consisting of
 Hammerhead, Inozyme, G-cleaver, DNazyme, Amberyze and Zinzyme. The
 nucleic acid molecules further comprise at least five ribose residues, at
 least ten 2'-O-methyl modifications, phosphorothioate linkages on at
 least three of the 5' terminal nucleotides and a 3' end modification of a
 3'-3', inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
 are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
 in the specification. The present sequence is that of a nucleic acid
 molecule of the invention

XX
 XX Sequence 17 BP; 2 A; 7 C; 6 G; 0 T; 2 U; 0 Other;

Query Match 78.9%; Score 15; DB 6; Length 17;
 Best Local Similarity 86.7%; Pred. No. 6.4e+02;
 Matches 13; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGT 15
 |||||:|||||:
 Db 3 CGGAAGCCCGCGU 17

RESULT 12

ADN36699/c

ID ADN36699 standard; DNA; 23 BP.

XX
 XX ADN36699;

XX
 XX 15-JUL-2004 (first entry)

XX
 XX West Nile virus detection-related PCR primer SeqID21.

XX
 XX hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; PCR; primer; ss.

XX
 XX West Nile virus.

XX
 XX WO2004036190-A2.

XX
 XX 29-APR-2004.

XX

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

PR 25-NOV-2002; 2002US-0429006P.

PR 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of
 bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Example 2; SEQ ID NO 21; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 detecting a nucleic acid, which is a probe sequence that comprises a
 target-complementary sequence of bases, and optionally one or more base
 sequences that are not complementary to the nucleic acid that is to be
 detected. The hybridisation assay probes and the kits are useful in
 detecting and amplifying a target nucleic acid sequence, for example
 flavivirus like West Nile virus, that may be present in a biological
 sample. West Nile virus (WNV) is an RNA virus that primarily infects
 birds and culex mosquitoes, with humans and horses serving as incidental
 hosts. Infection of humans can lead to meningitis or encephalitis. The
 invention may allow for accurate and efficient high throughput screening.
 The present sequence is that of a PCR primer which is related to the
 invention.

XX Sequence 23 BP; 8 A; 6 C; 7 G; 2 T; 0 U; 0 Other;

Query Match 78.9%; Score 15; DB 12; Length 23;
 Best Local Similarity 100.0%; Pred. No. 6.5e+02;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 5 ATGCCCGCGGTGTG 19
 |||||:|||||:
 Db 23 ATGCCCGCGGTGTG 9

RESULT 13

ADN36711/c

ID ADN36711 standard; DNA; 50 BP.

XX
 XX ADN36711;

XX 15-JUL-2004 (first entry)

XX West Nile virus detection-related oligonucleotide probe SeqID33.

XX hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.

XX West Nile virus.

XX Enterobacteria phage 77.

XX Key Location/Qualifiers

FT misc_feature 1..27
 FT /tag= a
 FT /note= "T7 promoter sequence"

FT misc_feature 28..50
 FT /tag= b
 FT /note= "WNV-complimentary sequence"

XX WO2004036190-A2.

XX 29-APR-2004.

XX

PF 10-OCT-2003; 2003WO-US033639.
 XX
 PR 16-OCT-2002; 2002US-0418891P.
 PR 25-NOV-2002; 2002US-0429006P.
 XX 24-FEB-2003; 2003US-0449810P.
 XX
 PA (GENP-) GEN-PROBE INC.
 XX
 PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX
 XX WPI; 2004-389590/36.
 DR
 XX
 XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 PT
 XX
 XX Disclosure; SEQ ID NO 33; 135pp; English.
 PS
 XX
 XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.
 XX
 SQ Sequence 50 BP; 19 A; 10 C; 12 G; 9 T; 0 U; 0 Other;
 Query Match 78.9%; Score 15; DB 12; Length 50;
 Best Local Similarity 100.0%; Pred. No. 6.7e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 5 ATGCCCCCGGTGGT 19
 DB 50 ATGCCCCCGGTGGT 36
 RESULT 14
 ADQ30683
 ID ADQ30683 standard; DNA; 19 BP.
 XX
 AC ADQ30683;
 XX
 DT 23-SEP-2004 (first entry)
 XX
 DE West Nile Virus oligonucleotide probe B.
 XX
 XX ss; probe; West Nile Virus; diagnosis.
 KW
 XX
 OS West Nile virus.
 OS
 PN WO2004055159-A2.
 PN
 PD 01-JUL-2004.
 XX
 PF 05-DEC-2003; 2003WO-US038750.
 XX
 PR 12-DEC-2002; 2002US-0432850P.
 PR 20-JUN-2003; 2003US-0480431P.
 XX
 XX (CHIR) CHIRON CORP.
 PA
 XX Shyamala V;
 PI
 XX WPI; 2004-488058/46.
 DR
 XX
 XX New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.

PT blood samples.
 XX
 PS Claim 1; SEQ ID NO 53; 56pp; English.
 XX
 CC The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to an oligonucleotide probe of the invention.
 XX
 SQ Sequence 19 BP; 2 A; 6 C; 7 G; 4 T; 0 U; 0 Other;
 Query Match 74.7%; Score 14.2; DB 12; Length 19;
 Best Local Similarity 84.2%; Pred. No. 1.7e+03;
 Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 CGGAATGCCCGCGTGGT 19
 DB 1 CGGTATGCCCGCGGATTG 19
 RESULT 15
 ADQ30679
 ID ADQ30679 standard; DNA; 19 BP.
 XX
 AC ADQ30679;
 XX
 DT 23-SEP-2004 (first entry)
 XX
 DE West Nile Virus capsid gene second probe.
 XX
 XX ss; probe; West Nile Virus; diagnosis.
 KW
 XX
 OS West Nile virus.
 OS
 PN WO2004055159-A2.
 PN
 PD 01-JUL-2004.
 XX
 PF 05-DEC-2003; 2003WO-US038750.
 XX
 PR 12-DEC-2002; 2002US-0432850P.
 PR 20-JUN-2003; 2003US-0480431P.
 XX
 XX (CHIR) CHIRON CORP.
 PA
 XX Shyamala V;
 PI
 XX WPI; 2004-488058/46.
 DR
 XX
 XX New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.
 PS Example 1; SEQ ID NO 49; 56pp; English.
 XX
 XX The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label

CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine (TAMRA), and 2',4',5',7'-tetrachloro-4,7-dichlorofluorescein (TET). The composition and methods are useful for accurately diagnosing West Nile virus infection or for capturing, detecting and quantitating West Nile virus in biological samples, particularly blood samples. This sequence corresponds to a probe to detect amplification of a fragment of the capsid gene of the WNV genome. The fragment is detected using the CC oligonucleotides of the invention.

XX SQ Sequence 19 BP; 2 A; 6 C; 7 G; 4 T; 0 U; 0 Other;
Query Match 74.7%; Score 14.2; DB 12; Length 19;
Best Local Similarity 84.2%; Pred. No. 1.7e+03;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGTG 19
Db 1 CGGTATGCCCGCGGATG 19

RESULT 16
ACN03482
ID ACN03482 standard; RNA; 17 BP.
XX ACN03482;
XX 22-APR-2004 (first entry)
XX WNV Zinzyne substrate SEQ ID NO 3485.
XX WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic; virucide; neuroprotective; antibacterial; replication; pancreatitis; encephalitis; myocarditis; meningitis; infection; hepatitis; liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNazyme; Amberzyme; Zinzyne; ss.
XX West Nile Virus.
XX WO200268637-A2.
XX 06-SEP-2002.
XX 19-OCT-2001; 2001WO-US048350.
XX 20-OCT-2000; 2000US-0242411P.
XX (RIBO-) RIBOZYME PHARM INC.
XX (BLAT/) BLATT L.
XX (MCSW/) MCSWIGGEN J A.
XX Blatt L, Mcswiggen JA;
XX WPI; 2002-706994/76.
XX New nucleic acid molecule that modulates replication of West Nile Virus (WNV), useful for treating a condition related to WNV infection e.g. pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX Claim 23; SEQ ID NO 3485; 495pp; English.

XX The invention relates to nucleic acid molecules that modulate replication of the West Nile Virus (WNV). The nucleic acid molecules are useful for treating a condition related to WNV infection e.g. pancreatitis, encephalitis, myocarditis, meningitis, neurologic infection, hepatitis, liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid molecule is selected from the group of ribozymes consisting of Hammerhead, Inozyme, G-cleaver, DNazyme, Amberzyme and Zinzyne. The nucleic acid molecules further comprise at least five ribose residues, at least ten 2'-O-methyl modifications, phosphorothioate linkages on at least three of the 5' terminal nucleotides and a 3' end modification of a 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080 are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given in the specification. The present sequence is that of a nucleic acid

CC molecule of the invention
XX SQ Sequence 17 BP; 0 A; 7 C; 5 G; 0 T; 5 U; 0 Other;
Query Match 73.7%; Score 14; DB 6; Length 17;
Best Local Similarity 71.4%; Pred. No. 2.1e+03;
Matches 10; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Qy 6 TGCCCCCGCGTGTG 19
Db 1 UGCCCCCGGUGUUG 14

RESULT 17
ACN12235/c
ID ACN12235 standard; RNA; 17 BP.
XX ACN12235;
XX 22-APR-2004 (first entry)
XX WNV minus strand Inozyme substrate SEQ ID NO 12238.
XX WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic; virucide; neuroprotective; antibacterial; replication; pancreatitis; encephalitis; myocarditis; meningitis; infection; hepatitis; liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNazyme; Amberzyme; Zinzyne; ss.
XX West Nile Virus.
XX WO200268637-A2.
XX 06-SEP-2002.
XX 19-OCT-2001; 2001WO-US048350.
XX 20-OCT-2000; 2000US-0242411P.
XX (RIBO-) RIBOZYME PHARM INC.
XX (BLAT/) BLATT L.
XX (MCSW/) MCSWIGGEN J A.
XX Blatt L, Mcswiggen JA;
XX WPI; 2002-706994/76.
XX New nucleic acid molecule that modulates replication of West Nile Virus (WNV), useful for treating a condition related to WNV infection e.g. pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX Claim 23; SEQ ID NO 12238; 495pp; English.

XX The invention relates to nucleic acid molecules that modulate replication of the West Nile Virus (WNV). The nucleic acid molecules are useful for treating a condition related to WNV infection e.g. pancreatitis, encephalitis, myocarditis, meningitis, neurologic infection, hepatitis, liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid molecule is selected from the group of ribozymes consisting of Hammerhead, Inozyme, G-cleaver, DNazyme, Amberzyme and Zinzyne. The nucleic acid molecules further comprise at least five ribose residues, at least ten 2'-O-methyl modifications, phosphorothioate linkages on at least three of the 5' terminal nucleotides and a 3' end modification of a 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080 are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given in the specification. The present sequence is that of a nucleic acid molecule of the invention

XX SQ Sequence 17 BP; 1 A; 6 C; 7 G; 0 T; 3 U; 0 Other;
Query Match 73.7%; Score 14; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. No. 2.1e+03;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```
QY      1 CGGAATGCCCGCG 14
Db      14 CGGAATGCCCGCG 1

RESULT 18
ADN36700/c
ID  ADN36700 standard; DNA; 23 BP.
XX
AC  ADN36700;
XX
DT  15-JUL-2004 (first entry)
XX
DE  West Nile virus detection-related PCR primer SeqID22.
XX
KW  hybridisation assay probe; nucleic acid detection;
KW  target-complementary sequence; flavivirus; West Nile virus; WNV;
KW  RNA virus; infection; meningitis; encephalitis;
KW  high throughput screening; PCR; primer; ss.
XX
OS  West Nile virus.
XX
FH  Key      Location/Qualifiers
FT  modified_base 12
FT  /*tag= a
FT  /*mod_base= i
XX
XX  WO2004036190-A2.
XX
XX  29-APR-2004.
XX
XX  10-OCT-2003; 2003WO-US033639.
XX
XX  16-OCT-2002; 2002US-0418891P.
XX  25-NOV-2002; 2002US-0429006P.
XX  24-FEB-2003; 2003US-0449810P.
XX
XX  (GENP-) GEN-PROBE INC.
XX
XX  Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX  WPI; 2004-389590/36.
XX
XX  New hybridization assay probe comprising target-complementary sequence of
XX  bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX  Example 2; SEQ ID NO 22; 135pp; English.
XX
XX  This invention relates to a novel hybridisation assay probe, for
XX  detecting a nucleic acid, which is a probe sequence that comprises a
XX  target-complementary sequence of bases, and optionally one or more base
XX  sequences that are not complementary to the nucleic acid that is to be
XX  detected. The hybridisation assay probes and the kits are useful in
XX  detecting and amplifying a target nucleic acid sequence, for example
XX  flavivirus like West Nile virus, that may be present in a biological
XX  sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX  birds and culex mosquitoes, with humans and horses serving as incidental
XX  hosts. Infection of humans can lead to meningitis or encephalitis. The
XX  invention may allow for accurate and efficient high throughput screening.
XX  The present sequence is that of a PCR primer which is related to the
XX  invention.
XX
XX  Sequence 23 BP; 8 A; 5 C; 7 G; 2 T; 0 U; 1 Other;
XX
XX  Query Match      73.7%; Score 14; DB 12; Length 23;
XX  Best Local Similarity 93.3%; Pred. No. 2.1e+03;
XX  Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
XX  QY      5 ATGCCCGCGGTG 19
XX      |||||
XX  Db      23 ATGCCCGCGGTG 9

RESULT 19
ADN36712/c
ID  ADN36712 standard; DNA; 50 BP.
XX
AC  ADN36712;
XX
DT  15-JUL-2004 (first entry)
XX
DE  West Nile virus detection-related oligonucleotide probe SeqID34.
XX
KW  hybridisation assay probe; nucleic acid detection;
KW  target-complementary sequence; flavivirus; West Nile virus; WNV;
KW  RNA virus; infection; meningitis; encephalitis;
KW  high throughput screening; probe; ss.
XX
OS  West Nile virus.
XX
FH  Key      Location/Qualifiers
FT  misc_feature 1..27
FT  /*tag= a
FT  /*note= "T7 promoter sequence"
FT  misc_feature 28..50
FT  /*tag= b
FT  /*note= "WNV-complementary sequence"
XX
XX  WO2004036190-A2.
XX
XX  29-APR-2004.
XX
XX  10-OCT-2003; 2003WO-US033639.
XX
XX  16-OCT-2002; 2002US-0418891P.
XX  25-NOV-2002; 2002US-0429006P.
XX  24-FEB-2003; 2003US-0449810P.
XX
XX  (GENP-) GEN-PROBE INC.
XX
XX  Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX  WPI; 2004-389590/36.
XX
XX  New hybridization assay probe comprising target-complementary sequence of
XX  bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX  Disclosure; SEQ ID NO 34; 135pp; English.
XX
XX  This invention relates to a novel hybridisation assay probe, for
XX  detecting a nucleic acid, which is a probe sequence that comprises a
XX  target-complementary sequence of bases, and optionally one or more base
XX  sequences that are not complementary to the nucleic acid that is to be
XX  detected. The hybridisation assay probes and the kits are useful in
XX  detecting and amplifying a target nucleic acid sequence, for example
XX  flavivirus like West Nile virus, that may be present in a biological
XX  sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX  birds and culex mosquitoes, with humans and horses serving as incidental
XX  hosts. Infection of humans can lead to meningitis or encephalitis. The
XX  invention may allow for accurate and efficient high throughput screening.
XX  The present sequence is that of an oligonucleotide probe which is related
XX  to the invention.
XX
XX  Sequence 50 BP; 19 A; 9 C; 12 G; 9 T; 0 U; 1 Other;
XX
XX  Query Match      73.7%; Score 14; DB 12; Length 50;
XX  Best Local Similarity 93.3%; Pred. No. 2.2e+03;
XX  Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
XX  QY      5 ATGCCCGCGGTG 19
XX      |||||
XX  Db      50 ATGCCCGCGGTG 36
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RESULT 20
ACN04726 standard; RNA; 17 BP.
AC ACN04726;
XX ACN04726;
XX 22-APR-2004 (first entry)
XX MNV DNAzyme substrate SEQ ID NO 4729.
XX DE
XX MNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
XX KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
XX KW encephalitis; myocarditis; meningitis; infection; hepatitis;
XX KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;
XX KW Amberzyme; Zinzyne; ss.
XX OS
XX West Nile Virus.
XX WO200268637-A2.
XX 06-SEP-2002.
XX 19-OCT-2001; 2001WO-US048350.
XX 20-OCT-2000; 2000US-0242411P.
XX (RIBO-) RIBOZYME PHARM INC.
XX (BLAT/) BLATT L.
XX (MCSW/) MCSWIGGEN J A.
XX Blatt L, Mcswiggen JA;
XX WPI; 2002-706994/76.
XX New nucleic acid molecule that modulates replication of West Nile Virus
XX (MNV), useful for treating a condition related to MNV infection e.g.
XX pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX Claim 23; SEQ ID NO 4729; 495pp; English.
XX The invention relates to nucleic acid molecules that modulate replication
XX of the West Nile Virus (MNV). The nucleic acid molecules are useful for
XX treating a condition related to MNV infection e.g. pancreatitis,
XX encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
XX liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
XX molecule is selected from the group of ribozymes consisting of
XX Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyne. The
XX nucleic acid molecules further comprise at least five ribose residues, at
XX least ten 2'-O-methyl modifications, phosphorothioate linkages on at
XX least three of the 5' terminal nucleotides and a 3' end modification of a
XX 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
XX are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
XX in the specification. The present sequence is that of a nucleic acid
XX molecule of the invention
XX SQ Sequence 17 BP; 4 A; 7 C; 5 G; 0 T; 1 U; 0 Other;
Query Match 68.4%; Score 13; DB 6; Length 17;
Best Local Similarity 92.3%; Pred. No. 6.8e+03;
Matches 12; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
Oy 1 CGGAATGCCCGC 13
Db 5 CGGAUGCCCGC 17
RESULT 21
ACN12234/c
ID ACN12234 standard; RNA; 17 BP.
XX ACN12234;
XX ACN12234;
XX 22-APR-2004 (first entry)
XX MNV minus strand Inozyme substrate SEQ ID NO 12237.
XX DE
XX MNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
XX KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
XX KW encephalitis; myocarditis; meningitis; infection; hepatitis;
XX KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;
XX KW Amberzyme; Zinzyne; ss.
XX OS
XX West Nile Virus.
XX WO200268637-A2.
XX 06-SEP-2002.
XX 19-OCT-2001; 2001WO-US048350.
XX 20-OCT-2000; 2000US-0242411P.
XX (RIBO-) RIBOZYME PHARM INC.
XX (BLAT/) BLATT L.
XX (MCSW/) MCSWIGGEN J A.
XX Blatt L, Mcswiggen JA;
XX WPI; 2002-706994/76.
XX New nucleic acid molecule that modulates replication of West Nile Virus
XX (MNV), useful for treating a condition related to MNV infection e.g.
XX pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX Claim 23; SEQ ID NO 12237; 495pp; English.
XX The invention relates to nucleic acid molecules that modulate replication
XX of the West Nile Virus (MNV). The nucleic acid molecules are useful for
XX treating a condition related to MNV infection e.g. pancreatitis,
XX encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
XX liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
XX molecule is selected from the group of ribozymes consisting of
XX Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyne. The
XX nucleic acid molecules further comprise at least five ribose residues, at
XX least ten 2'-O-methyl modifications, phosphorothioate linkages on at
XX least three of the 5' terminal nucleotides and a 3' end modification of a
XX 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
XX are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
XX in the specification. The present sequence is that of a nucleic acid
XX molecule of the invention
XX SQ Sequence 17 BP; 5 A; 5 C; 7 G; 0 T; 0 U; 0 Other;
Query Match 68.4%; Score 13; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. No. 6.8e+03;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 7 GCCCGCGCTGTG 19
Db 17 GCCCGCGCTGTG 5
RESULT 22
AAA49102
ID AAA49102 standard; DNA; 24 BP.
XX AAA49102;
XX AAA49102;
XX 16-NOV-2000 (first entry)
XX Forward primer 650-E-S used to synthesize snut 650-720-EcoRI.
XX HIV; human immunodeficiency virus; vaccine; AIDS; PCR primer; snut;
XX KW silent nucleotide substitution; ss.
XX OS Synthetic.
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XX PN WO200029561-A2.
XX PD 25-MAY-2000.
XX PF 27-MAR-2000; 2000WO-DK000144.
XX PR 29-MAR-1999; 99DK-00000427.
XX PR 09-APR-1999; 99US-0128558P.
XX PA (STAT-) STATENS SERUM INST.
XX PI Fomsgaard A;
XX PT WPI; 2000-387778/33.
XX PT Producing nucleotide sequence construct with optimized codons for human
XX PT immunodeficiency virus (HIV) genetic vaccine involves obtaining a first
XX PT nucleotide sequence from a HIV patient, redesigning and assembling it
XX PT with snuts.
XX PS Example 3; Page 30; 150pp; English.
XX CC The present invention relates to a nucleotide construct with optimised
XX CC codons for use as a human immunodeficiency virus (HIV) DNA vaccine. The
XX CC construct uses codons from highly expressed mammalian proteins to code
XX CC for each derivative of an early, primary HIV envelope gene. The first
XX CC stage in the production of the construct was the cloning of an HIV
XX CC envelope gene. A nucleotide sequence encoding this gene was then created
XX CC using codons from highly expressed mammalian genes. The present sequence
XX CC is a PCR primer that was used to clone one of the snuts (AAA49060-AA9079)
XX CC that were created by redesigning this nucleotide construct so that
XX CC restriction enzyme sites surrounded functional regions of the sequence.
XX CC The snuts were then assembled into pieces (AAA49080-AA9092). Each
XX CC derivative of the envelope gene (AAA49093-AA9097) was then built using
XX CC the pieces. The HIV DNA vaccine may be used as a prophylactic vaccine and
XX CC as a therapeutic vaccine in HIV infected patients
XX SQ Sequence 24 BP; 4 A; 8 C; 8 G; 4 T; 0 U; 0 Other;

Query Match 66.3%; Score 12.6; DB 3; Length 24;
Best Local Similarity 78.9%; Pred. No. 1.1e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CGGAATGCCCGCGTGTG 19
Db ||||| ||||| ||||| ||||| |||||
2 CGGAATGCCCGCGTGTG 20

RESULT 23
AAI30978/c
ID AAI30978 standard; DNA; 31 BP.
XX AC AAI30978;
XX DT 04-NOV-2004 (revised)
XX DT 18-OCT-2001 (first entry)
XX DE Human single nucleotide polymorphism (SNP) HOXB3.
XX KW Human; resequence; genotype; disease; forensic; paternity testing;
XX KW single nucleotide polymorphism; SNP; ss.
XX OS Homo sapiens.
XX PH Key Location/Qualifiers
XX FT variation 16
XX FT /*tag= a
XX FT /standard_name= "single nucleotide polymorphism"
XX PN WO200166800-A2.
XX PD 13-SEP-2001.

XX PF 07-MAR-2001; 2001WO-US007268.
XX PR 07-MAR-2000; 2000US-0187510P.
XX PR 22-MAY-2000; 2000US-0206129P.
XX PA (WHED ) WHITEHEAD INST BIOMEDICAL RES.
XX PI Cargill M, Ireland JS, Lander ES;
XX DR WPI; 2001-522952/57.
XX CC Nucleic acid molecules from the human genome which include polymorphic
XX CC sites, useful in methods for predicting the presence, absence or severity
XX CC of a particular phenotype or disorder (e.g. diabetes) associated with a
XX CC particular genotype.
XX PS Claim 1; Page 120; 145pp; English.
XX CC The invention relates to the identification of nucleic acid molecules
XX CC (AAI29513-AAI31314) from the human genome which include polymorphic sites
XX CC of individuals were resequenced and single nucleotide polymorphisms
XX CC (SNPs) in these genes discovered. The method is useful for predicting the
XX CC presence, absence or severity of a particular phenotype or disorder (e.g.
XX CC diabetes) associated with a particular genotype. The nucleic acids
XX CC containing the polymorphic sites may be useful in forensics and paternity
XX CC testing
XX CC Revised record issued on 04-NOV-2004 : Correction to Feature Table Key
XX SQ Sequence 31 BP; 5 A; 9 C; 12 G; 5 T; 0 U; 0 Other;

Query Match 66.3%; Score 12.6; DB 4; Length 31;
Best Local Similarity 78.9%; Pred. No. 1.1e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CGGAATGCCCGCGTGTG 19
Db ||||| ||||| ||||| ||||| |||||
19 CGGAATGCCCGCGTGTG 1

RESULT 24
AAI50260/c
ID AAI50260 standard; DNA; 45 BP.
XX AC AAI50260;
XX DT 13-FEB-2003 (first entry)
XX DE Antigenic chimeric protein coding sequence fragment #2.
XX KW Antigen; oligomerisation domain; oligomeric protein complex; virucide;
XX KW vaccine; PACGCN4nas; influenza; chimera; immunostimulant; gene; ds.
XX OS Synthetic.
XX PH Key Location/Qualifiers
XX FT CDS complement(4. .45)
XX FT /*tag= a
XX FT /product= "Antigenic chimeric protein fragment"
XX PN WO200274795-A2.
XX PD 26-SEP-2002.
XX PF 18-JAN-2002; 2002WO-EP000628.
XX PR 18-JAN-2001; 2001EP-00200193.
XX PA (VLAA-) VLAAms INTERUNIVERSITAIR INST BIOTECHNOG.
XX PI De Fillette M, Deroo TW, Fiers W, Maras M, Min Jou WA;

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XX WPI; 2003-058336/05.
DR P-PSDB; AAO19602.
XX
XX New chimeric protein comprising an antigen derived from a naturally
PT occurring oligomeric protein complex, and an oligomerization domain,
PT useful in preparing a vaccine against influenza.
XX
XX Example 1; Fig 1; 53pp; English.
XX
XX The present invention provides chimeric proteins each comprising an
CC antigen derived from a naturally occurring oligomeric protein complex and
CC an oligomerization domain. These can be used in the preparation of
CC vaccines, particularly against influenza. The present sequence encodes a
CC fragment of a chimeric protein of the invention
XX
XX Sequence 45 BP; 11 A; 11 C; 12 G; 11 T; 0 U; 0 Other;
SQ
Query Match 66.3%; Score 12.6; DB 8; Length 45;
Best Local Similarity 78.9%; Pred. No. 1.1e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 1 CGGAATGCCCGCGGTGTG 19
Db 37 CGGACTACCCCGCTGTAG 19
RESULT 25
ABL99256/C
ID ABL99256 standard; DNA; 54 BP.
XX
XX ABL99256;
AC
XX
XX 27-JUN-2002 (first entry)
XX
XX Green/red click beetle luciferase preparation oligo SEQ ID NO:234.
XX
XX Luciferase; synthetic nucleic acid; transcriptional characteristic;
KW transcription; codon usage; PCR; primer; ss.
XX
XX Coleoptera.
OS
XX Synthetic.
XX
XX WO200216944-A2.
XX
XX 28-FEB-2002.
XX
XX 24-AUG-2001; 2001WO-US026566.
XX
XX 24-AUG-2000; 2000US-00645706.
XX
XX (PROM-) PROMEGA CORP.
XX
XX Wood KV, Wood MG, Zhuang Y, Paguio A;
XX
XX WPI; 2002-304140/34.
XX
XX Preparing a synthetic nucleic acid molecule with reduced inappropriate
PT transcriptional characteristics when expressed in a cell, for e.g making
PT fusion proteins, by altering a wild type or another synthetic nucleic
PT acid sequence.
XX
XX Example 1; Fig 6; 294pp; English.
XX
XX The present invention relates to the preparation of synthetic nucleic
CC acid molecules which have altered transcriptional regulatory sequences
CC compared to the wild-type. These sequences are then transcribed with less
CC frequency compared to the wild-type. In particular, the invention relates
CC to altered luciferase sequences. This can be used to detect weak promoter
CC activity, to express fusion proteins, to detect and/or measure levels of
CC gene expression, subcellular localisation or targeting, in life science
CC research, agro genetics, gene therapy, developmental science and
CC pharmaceutical development. The present sequence is an oligonucleotide
```

```
CC described in the exemplification of the invention
XX
XX Sequence 54 BP; 16 A; 11 C; 16 G; 11 T; 0 U; 0 Other;
SQ
Query Match 66.3%; Score 12.6; DB 6; Length 54;
Best Local Similarity 78.9%; Pred. No. 1.2e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 1 CGGAATGCCCGCGGTGTG 19
Db 19 CGGAATGCCCAAGCTTTTG 1
RESULT 26
AAA76109/C
ID AAA76109 standard; DNA; 60 BP.
XX
XX AAA76109;
AC
XX
XX 11-DEC-2000 (first entry)
XX
XX Aspergillus awamori glucoamylase mutagenic primer #10.
DE
XX
XX Glucoamylase; enzyme; carboxydase; glucose;
KW 1,4-alpha-D-glucan glucohydrolase; mutagenic primer; ss.
XX
XX Aspergillus awamori.
OS
XX
XX WO2000043504-A1.
XX
XX 27-JUL-2000.
XX
XX 10-JAN-2000; 2000WO-US000532.
XX
XX 22-JAN-1999; 99US-00236063.
XX
XX (IOWA) UNIV IOWA STATE RES FOUND INC.
XX
XX Allen MJ, Fang T, Li Y, Liu H, Chen H, Coutinho P, Honzatko R;
PI Ford C;
XX
XX WPI; 2000-514725/46.
XX
XX Fungal glucoamylase for selective production of glucose rather than alpha
PT -1,6 linked disaccharide isomaltose, has mutation pair Asn20Cys coupled
PT with Ala27Cys forming disulfide bond between the two stabilizing members.
XX
XX Example 5; Page 48; 160pp; English.
XX
XX The present invention relates to mutant glucoamylases (1,4-alpha-D-glucan
CC glucohydrolase; E.C. 3.2.1.3), which have increased thermostability,
CC increased pH optimum and reduced isomaltose formation. Glucoamylase is a
CC carboxydase, and cleaves D-glucose from the nonreducing ends of
CC maltooligosaccharides, attacking alpha-(1,4)-, and alpha-(1,6)-glucosidic
CC bonds. The mutant proteins (see AAB15178-B15184) are useful for the
CC selective production of glucose rather than alpha-1,6 linked disaccharide
CC isomaltose. The present sequence is a mutagenic primer used in the
CC generation of the mutant glucoamylases of the present invention
XX
XX Sequence 60 BP; 14 A; 20 C; 17 G; 9 T; 0 U; 0 Other;
SQ
Query Match 66.3%; Score 12.6; DB 3; Length 60;
Best Local Similarity 78.9%; Pred. No. 1.2e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 1 CGGAATGCCCGCGGTGTG 19
Db 48 CGGTTGCCCTGCGAGTTG 30
RESULT 27
AAA76110/C
ID AAA76110 standard; DNA; 60 BP.
```

```

XX AAA76110;
AC
XX
XX 11-DEC-2000 (first entry)
DT
XX
XX Aspergillus awamori glucoamylase mutagenic primer #11.
DE
XX Glucoamylase; enzyme; carbohydrase; glucose;
KW
KW 1,4-alpha-D-glucan glucohydrolase; mutagenic primer; ss.
XX
XX Aspergillus awamori.
OS
XX WO200043504-A1.
XX
XX 27-JUL-2000.
PD
XX
XX 10-JAN-2000; 2000WO-US000532.
PF
XX 22-JAN-1999; 99US-00236063.
XX
XX (IOWA ) UNIV IOWA STATE RES FOUND INC.
PA
XX Allen MJ, Fang T, Li Y, Liu H, Chen H, Coutinho P, Honzatko R;
PI Ford C;
PI
XX WPI; 2000-514725/46.
XX
XX Fungal glucoamylase for selective production of glucose rather than alpha
PT -1,6 linked disaccharide isomaltose, has mutation pair Asn20Cys coupled
PT with Ala27Cys forming disulfide bond between the two stabilizing members.
PT
XX Example 5; Page 48; 160pp; English.
PS
XX The present invention relates to mutant glucoamylases (1,4-alpha-D-glucan
CC glucosylase; E.C. 3.2.1.3), which have increased thermostability,
CC increased pH optimum and reduced isomaltose formation. Glucoamylase is a
CC carbohydrase, and cleaves D-glucose from the nonreducing ends of
CC maltooligosaccharides, attacking alpha-(1,4)-, and alpha-(1,6)-glucosidic
CC bonds. The mutant proteins (see AAB15178-B15184) are useful for the
CC selective production of glucose rather than alpha-1,6 linked disaccharide
CC isomaltose. The present sequence is a mutagenic primer used in the
CC generation of the mutant glucoamylases of the present invention
XX
SQ Sequence 60 BP; 14 A; 20 C; 17 G; 9 T; 0 U; 0 Other;

Query Match 66.3%; Score 12.6; DB 3; Length 60;
Best Local Similarity 78.9%; Pred. No. 1.2e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CGGATGCCCGCGGTGG 19
    ||| ||| ||| ||| |||
DB 48 CGGGTGGCCCTGCGAGTTG 30

RESULT 28
ACH55782/c
ID ACH55782 standard; DNA; 25 BP.
XX
AC ACH55782;
XX
XX 16-OCT-2003 (first entry)
DT
XX
XX DNA target sequence #4918 useful in array for genetic analyses.
DE
XX
XX Gene expression analysis; array; hybridisation; genetic variation;
KW tag-labelled compound; gene family; in situ hybridisation;
KW library screening; Southern hybridisation; northern hybridisation;
KW dot-blot hybridisation; gene sequence; mutation detection;
KW target sequence; probe; PCR; primer; ss.
XX
XX Unidentified.
OS
XX US2003082596-A1.
PN

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XX 01-MAY-2003.
PD
XX
XX 08-AUG-2002; 2002US-00215112.
PF
XX
XX 08-AUG-2001; 2001US-0311040P.
PR
XX (MITT/) MITTMANN M.
PA
XX Mittmann M;
XX
XX WPI; 2003-576608/54.
XX
XX New probe array useful e.g. for monitoring gene expression levels, for
PT analyzing genetic variations, or for hybridizing tag-labeled compounds,
PT comprises multiple nucleic acid probes.
PT
XX Claim 1; SEQ ID NO 4918; 9pp; English.
XX
XX The present invention relates to nucleic acid sequences that are
CC complementary to particular genes, and can be used as probes for a
CC variety of analyses such as gene expression analysis. Each probe
CC comprises 9 or more consecutive nucleotides from at least one of 14936
CC nucleotide sequences defined in the patent, or their perfect sense match,
CC sense mismatch, antisense match or antisense mismatch oligonucleotides.
CC The probes may be used in an array comprising at least 10 distinct
CC nucleic acid probes. The array is useful in monitoring gene expression
CC levels by hybridisation to a DNA library, in analysing genetic
CC variations, and in hybridising tag-labelled compounds. The probes are
CC useful for identifying family members of a gene. The probes are also
CC useful in situ hybridisations, in screening cDNA or genomic libraries
CC (or derived subclones) for additional clones containing segments of DNA
CC that have been previously isolated and sequenced in Southern, northern,
CC or dot-blot hybridisation of genomic DNA to identify or detect the
CC sequence of any gene or detect specific mutations in any gene, and in
CC mapping the 5' termini of mRNA molecules by primer extensions. The
CC nucleic acid sequences of the invention are also useful as PCR primers.
CC The invention provides a large collection of nucleic acid sequences
CC complementary to particular genes with a wide range of analytical uses.
CC ACH50865-ACH5260 represent the target sequences of the invention. Note:
CC The sequence data for this patent was obtained in electronic format
CC directly from the USPTO web site at seqdata.uspto.gov/paipsDIDentry.html
XX
SQ Sequence 25 BP; 6 A; 9 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 65.3%; Score 12.4; DB 9; Length 25;
Best Local Similarity 92.9%; Pred. No. 1.4e+04;
Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 TGCCCGCGGTGG 19
    ||| ||| ||| ||| |||
DB 22 TGCACCGCGGTGG 9

RESULT 29
ADN36841
ID ADN36841 standard; RNA; 25 BP.
XX
AC ADN36841;
XX
XX 15-JUL-2004 (first entry)
DT
XX
XX West Nile virus detection-related oligonucleotide probe SeqID163.
DE
XX
XX hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX
XX West Nile virus.
OS
XX
XX Key Location/Qualifiers
FH modified_base 1..25
FT

```

```
FT      /*tag= a
FT      /mod_base= OTHER
FT      /note= "OTHER= 2'-methoxyethoxy (2'-MOE) nucleotides"
XX
XX      WO2004036190-A2.
XX
XX      29-APR-2004.
XX
XX      10-OCT-2003; 2003WO-US033639.
XX
XX      16-OCT-2002; 2002US-0418891P.
XX
XX      25-NOV-2002; 2002US-0429006P.
XX
XX      24-FEB-2003; 2003US-0449810P.
XX
XX      (GENP-) GEN-PROBE INC.
XX
XX      Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX
XX      WPI; 2004-389590/36.
XX
XX      New hybridization assay probe comprising target-complementary sequence of
XX      bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX      Example 7; SEQ ID NO 163; 135pp; English.
XX
XX      This invention relates to a novel hybridisation assay probe, for
XX      detecting a nucleic acid, which is a probe sequence that comprises a
XX      target-complementary sequence of bases, and optionally one or more base
XX      sequences that are not complementary to the nucleic acid that is to be
XX      detected. The hybridisation assay probes and the kits are useful in
XX      detecting and amplifying a target nucleic acid sequence, for example
XX      flavivirus like West Nile virus, that may be present in a biological
XX      sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX      birds and culex mosquitoes, with humans and horses serving as incidental
XX      hosts. Infection of humans can lead to meningitis or encephalitis. The
XX      invention may allow for accurate and efficient high throughput screening.
XX      The present sequence is that of an oligonucleotide probe which is related
XX      to the invention.
XX
XX      Sequence 25 BP; 6 A; 9 C; 8 G; 0 T; 2 U; 0 Other;
XX
XX      Query Match      65.3%; Score 12.4; DB 12; Length 25;
XX      Best Local Similarity 85.7%; Pred. No. 1.4e+04;
XX      Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
XX
XX      Qy      1 CGGAATGCCCGCG 14
XX      |||||:|||||
XX      Db      11 CGGAATGCCCGCG 24
XX
XX      RESULT 30
XX      AAZ27629
XX      ID      AAZ27629 standard; DNA; 24 BP.
XX
XX      AC      AAZ27629;
XX
XX      DT      20-DEC-1999 (first entry)
XX
XX      DE      PCR primer for GUS gene.
XX
XX      KW      Extracellular compartment modification; floral cell; self-compatibility;
XX      pollen-pistil interaction; self-incompatibility; insect growth control;
XX      PCR primer; GUS gene; ss.
XX
XX      OS      Synthetic.
XX      OS      Nicotiana tabacum.
XX
XX      PN      WO9949063-A1.
XX
XX      PD      30-SEP-1999.
XX
XX      PF      19-MAR-1999; 99WO-CA000237.
XX
XX
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```
PR      20-MAR-1998; 98US-0078728P.
XX
XX      PA      (MIAC ) CANADA MIN AGRIC & AGRI-FOOD CANADA.
XX
XX      PI      Robert LS, Gleddie S;
XX
XX      DR      WPI; 1999-591104/50.
XX
XX      Protein expression in floral cells for peptide display, mediating plant
XX      sterility, and modifying pollen-pistil interactions.
XX
XX      Example 12; Page 50; 113pp; English.
XX
XX      This sequence represents a PCR primer for the Nicotiana tabacum GUS gene.
XX      The invention relates to a method for modifying the extracellular
XX      compartment of a floral cell of a plant, that comprises expressing a
XX      construct comprising a gene of interest encoding a protein, fusion
XX      protein or peptide, or a fragment of them, which is capable of modifying
XX      the composition of the extracellular compartment of the floral cell and
XX      altering either the function, use or development of the floral cell or
XX      modifying the interaction of the floral cell with other cells, within an
XX      anther or pistil cell. The method is used to modify pollen-pistil
XX      interaction or function, which mediates, produces or prevents self-
XX      compatibility, self-incompatibility, out- or in-crossing or combinations
XX      of these. The method is also used for localizing proteins on the surface
XX      of pollen for the purpose of peptide display. The protein localized on
XX      the surface of the pollen may be an antibody or antigen or is a protein
XX      that is effective in controlling insect growth, behaviour, feeding,
XX      development or reproduction
XX
XX      Sequence 24 BP; 4 A; 7 C; 6 G; 7 T; 0 U; 0 Other;
XX
XX      Query Match      64.2%; Score 12.2; DB 2; Length 24;
XX      Best Local Similarity 82.4%; Pred. No. 1.8e+04;
XX      Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
XX
XX      Qy      2 GGAATGCCCGCGTGT 18
XX      |||||:|||||
XX      Db      1 GGAATTCACGCGTCTT 17
XX
XX      Search completed: September 9, 2005, 21:43:06
XX      Job time : 271 secs
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 9, 2005, 11:24:12 ; Search time 1774 Seconds
(without alignments)
407.678 Million cell updates/sec

Title: US-10-729-421-52
Perfect score: 19
Sequence: 1 cggatgccccgcgtgttg 19

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 241816

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 150 summaries

Database : EST:
1: gb_est1:*
2: gb_est2:*
3: gb_hic:*
4: gb_est3:*
5: gb_est4:*
6: gb_est5:*
7: gb_est6:*
8: gb_gss1:*
9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|---------------------|
| C 1 | 13.8 | 72.6 | 55 | 1 | AA776363 ahild05.s |
| C 2 | 13.4 | 70.5 | 39 | 1 | AV836760 AV836760 |
| C 3 | 13.2 | 69.5 | 54 | 1 | AV841787 AV841787 |
| C 4 | 13 | 68.4 | 30 | 4 | BM396904 BM396904 |
| C 5 | 12.8 | 67.4 | 54 | 2 | AV951548 AV951548 |
| C 6 | 12.6 | 66.3 | 36 | 4 | BG717269 BG717269 |
| C 7 | 12.6 | 66.3 | 37 | 4 | BG722105 BG722105 |
| C 8 | 12.6 | 66.3 | 40 | 1 | AI745090 t220d07.x |
| C 9 | 12.6 | 66.3 | 58 | 9 | BX892307 BX892307 |
| C 10 | 12.4 | 65.3 | 30 | 4 | BM398127 BM398127 |
| C 11 | 12.4 | 65.3 | 52 | 7 | CF325296 CF325296 |
| C 12 | 12 | 63.2 | 22 | 4 | BM398778 BM398778 |
| C 13 | 12 | 63.2 | 34 | 1 | AI646800 AI646800 |
| C 14 | 12 | 63.2 | 58 | 9 | CR078210 CR078210 |
| C 15 | 11.8 | 62.1 | 38 | 9 | TA274G03Q TA274G03Q |
| C 16 | 11.8 | 62.1 | 52 | 1 | AI958756 AI958756 |
| C 17 | 11.8 | 62.1 | 55 | 9 | CR770268 CR770268 |
| C 18 | 11.6 | 61.1 | 26 | 8 | AZ333170 AZ333170 |
| C 19 | 11.6 | 61.1 | 48 | 7 | CO781458 CO781458 |
| C 20 | 11.6 | 61.1 | 51 | 8 | BZ662469 BZ662469 |
| C 21 | 11.4 | 60.0 | 27 | 4 | BM398263 BM398263 |
| C 22 | 11.4 | 60.0 | 30 | 4 | BM396162 BM396162 |
| C 23 | 11.4 | 60.0 | 34 | 4 | BM400993 BM400993 |
| C 24 | 11.4 | 60.0 | 50 | 1 | AU105629 AU105629 |

| | | | | | | | |
|----------|-----------|----|---|-----------|------|------|------|
| AZ769819 | 1M0570E20 | 27 | 8 | AZ769819 | 58.9 | 11.2 | 25 |
| AI001661 | EST0243 T | 31 | 1 | AI001661 | 58.9 | 11.2 | 26 |
| BM400565 | 5009-0-75 | 31 | 4 | BM400565 | 58.9 | 11.2 | C 27 |
| BM398283 | 5009-0-43 | 33 | 4 | BM398283 | 58.9 | 11.2 | C 28 |
| AU009955 | AU009955 | 39 | 1 | AU009955 | 58.9 | 11.2 | C 29 |
| AI544749 | fb66g01.x | 46 | 1 | AI544749 | 58.9 | 11.2 | C 30 |
| BM400051 | 5009-0-65 | 46 | 1 | BM400051 | 58.9 | 11.2 | C 31 |
| AI685633 | tt89c09.x | 49 | 1 | AI685633 | 58.9 | 11.2 | C 32 |
| AU104742 | AU104742 | 50 | 1 | AU104742 | 58.9 | 11.2 | C 33 |
| AU105257 | AU105257 | 50 | 1 | AU105257 | 58.9 | 11.2 | C 34 |
| AA795813 | vv26h01.f | 56 | 1 | AA795813 | 58.9 | 11.2 | C 35 |
| BG730068 | de09d07.y | 57 | 1 | BG730068 | 58.9 | 11.2 | C 36 |
| CB173165 | OR_2027C1 | 57 | 6 | CB173165 | 58.9 | 11.2 | C 37 |
| AU472147 | T. brucei | 57 | 9 | TA149B12P | 58.9 | 11.2 | C 38 |
| CN924300 | 000414AEL | 60 | 7 | CN924300 | 58.9 | 11.2 | C 39 |
| AZ290346 | 1006019E0 | 60 | 8 | AZ290346 | 58.9 | 11.2 | C 40 |
| AI197517 | Tetraodon | 60 | 9 | CNS02HEC | 58.9 | 11.2 | C 41 |
| BG503651 | 602549510 | 61 | 4 | BG503651 | 57.9 | 11 | C 42 |
| CC199749 | XH279 Bay | 45 | 8 | CC199749 | 57.9 | 11 | C 43 |
| AI813520 | wj83D06.x | 49 | 1 | AI813520 | 57.9 | 11 | C 44 |
| AU106802 | AU106802 | 50 | 1 | AU106802 | 57.9 | 11 | C 45 |
| CB751759 | TGESTzyh5 | 51 | 6 | CB751759 | 57.9 | 11 | C 46 |
| AI652258 | wb28c09.x | 54 | 1 | AI652258 | 57.9 | 11 | C 47 |
| CF889771 | TcTR-574 | 54 | 7 | CF889771 | 57.9 | 11 | C 48 |
| AI903643 | QV-BT032- | 55 | 1 | AI903643 | 57.9 | 11 | C 49 |
| CR073588 | Forward s | 56 | 9 | CR073588 | 57.9 | 11 | C 50 |
| AI326751 | mq64h10.y | 57 | 1 | AI326751 | 57.9 | 11 | C 51 |
| CB369437 | TGESTzyg8 | 57 | 6 | CB369437 | 57.9 | 11 | C 52 |
| CB382289 | TGESTzyg7 | 57 | 6 | CB382289 | 57.9 | 11 | C 53 |
| CB382303 | TGESTzyg7 | 57 | 6 | CB382303 | 57.9 | 11 | C 54 |
| CB382546 | TGESTzyg8 | 57 | 6 | CB382546 | 57.9 | 11 | C 55 |
| CB382629 | TGESTzyg8 | 57 | 6 | CB382629 | 57.9 | 11 | C 56 |
| CB382752 | TGESTzyg9 | 57 | 6 | CB382752 | 57.9 | 11 | C 57 |
| CB382790 | TGESTzyg9 | 57 | 6 | CB382790 | 57.9 | 11 | C 58 |
| CB383084 | TGESTzyh4 | 57 | 6 | CB383084 | 57.9 | 11 | C 59 |
| CB383746 | TGESTzyg8 | 57 | 6 | CB383746 | 57.9 | 11 | C 60 |
| CB384264 | TGESTzyh5 | 57 | 6 | CB384264 | 57.9 | 11 | C 61 |
| CB412068 | TGESTzyh4 | 57 | 6 | CB412068 | 57.9 | 11 | C 62 |
| CB751697 | TGESTzyh5 | 57 | 6 | CB751697 | 57.9 | 11 | C 63 |
| CB752652 | TGESTzyh7 | 57 | 6 | CB752652 | 57.9 | 11 | C 64 |
| CB752662 | TGESTzyh7 | 57 | 6 | CB752662 | 57.9 | 11 | C 65 |
| CB752949 | TGESTzyh7 | 57 | 6 | CB752949 | 57.9 | 11 | C 66 |
| CB753824 | TGESTzyh9 | 57 | 6 | CB753824 | 57.9 | 11 | C 67 |
| CB754684 | TGESTzyh9 | 57 | 6 | CB754684 | 57.9 | 11 | C 68 |
| CR002115 | Forward s | 57 | 9 | CR002115 | 57.9 | 11 | C 69 |
| AI365545 | q208g12.x | 58 | 1 | AI365545 | 57.9 | 11 | C 70 |
| AJ747306 | AJ747306 | 58 | 1 | AJ747306 | 57.9 | 11 | C 71 |
| CB368700 | TGESTzyh1 | 58 | 6 | CB368700 | 57.9 | 11 | C 72 |
| CB369360 | TGESTzyg8 | 58 | 6 | CB369360 | 57.9 | 11 | C 73 |
| CB370433 | TGESTzyh1 | 58 | 6 | CB370433 | 57.9 | 11 | C 74 |
| CB383674 | TGESTzyg8 | 58 | 6 | CB383674 | 57.9 | 11 | C 75 |
| CB384012 | TGESTzyh4 | 58 | 6 | CB384012 | 57.9 | 11 | C 76 |
| CB754160 | TGESTzyh8 | 58 | 6 | CB754160 | 57.9 | 11 | C 77 |
| CB754316 | TGESTzyh9 | 58 | 6 | CB754316 | 57.9 | 11 | C 78 |
| CB755076 | TGESTzyh0 | 58 | 6 | CB755076 | 57.9 | 11 | C 79 |
| CB368397 | TGESTzyh0 | 59 | 6 | CB368397 | 57.9 | 11 | C 80 |
| CB368487 | TGESTzyh0 | 59 | 6 | CB368487 | 57.9 | 11 | C 81 |
| CB368662 | TGESTzyh0 | 59 | 6 | CB368662 | 57.9 | 11 | C 82 |
| CB369243 | TGESTzyg6 | 59 | 6 | CB369243 | 57.9 | 11 | C 83 |
| CB369683 | TGESTzyg7 | 59 | 6 | CB369683 | 57.9 | 11 | C 84 |
| CB369914 | TGESTzyh2 | 59 | 6 | CB369914 | 57.9 | 11 | C 85 |
| CB382011 | TGESTzyh3 | 59 | 6 | CB382011 | 57.9 | 11 | C 86 |
| CB382589 | TGESTzyg8 | 59 | 6 | CB382589 | 57.9 | 11 | C 87 |
| CB383643 | TGESTzyg8 | 59 | 6 | CB383643 | 57.9 | 11 | C 88 |
| CB383844 | TGESTzyg9 | 59 | 6 | CB383844 | 57.9 | 11 | C 89 |
| CB411517 | TGESTzyh3 | 59 | 6 | CB411517 | 57.9 | 11 | C 90 |
| CB751723 | TGESTzyh5 | 59 | 6 | CB751723 | 57.9 | 11 | C 91 |
| CB753658 | TGESTzyh9 | 59 | 6 | CB753658 | 57.9 | 11 | C 92 |
| CB411539 | TGESTzyg8 | 60 | 6 | CB411539 | 57.9 | 11 | C 93 |
| CB753796 | TGESTzyh9 | 60 | 6 | CB753796 | 57.9 | 11 | C 94 |
| BM397206 | 5009-0-3- | 23 | 4 | BM397206 | 56.8 | 10.8 | C 95 |
| AJ596200 | ArabiDops | 27 | 9 | AJ596200 | 56.8 | 10.8 | C 96 |
| AJ596226 | ArabiDops | 27 | 9 | AJ596226 | 56.8 | 10.8 | C 97 |

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c 98 10.8 56.8 40 1 A1491937
c 99 10.8 56.8 41 7 R19040
c 100 10.8 56.8 42 8 C181914
c 101 10.8 56.8 43 8 C181914
c 102 10.8 56.8 46 1 A182146
c 103 10.8 56.8 48 4 B1730786
c 104 10.8 56.8 48 9 CG721564
c 105 10.8 56.8 50 1 A1013160
c 106 10.8 56.8 50 1 A1063373
c 107 10.8 56.8 50 1 A106380
c 108 10.8 56.8 50 1 A107950
c 109 10.8 56.8 51 2 AV962960
c 110 10.8 56.8 52 7 CF298391
c 111 10.8 56.8 52 7 CF298391
c 112 10.8 56.8 59 2 AV966402
c 113 10.6 55.8 28 1 A937308
c 114 10.6 55.8 34 1 A288354
c 115 10.6 55.8 37 1 A1815205
c 116 10.6 55.8 42 9 CL520536
c 117 10.6 55.8 43 1 AA948558
c 118 10.6 55.8 43 1 AA221690
c 119 10.6 55.8 43 1 AL643698
c 120 10.6 55.8 49 8 CC178743
c 121 10.6 55.8 50 1 A10102246
c 122 10.6 55.8 50 1 A10102247
c 123 10.6 55.8 50 1 A10102248
c 124 10.6 55.8 50 1 A10102249
c 125 10.6 55.8 50 1 A10102250
c 126 10.6 55.8 50 1 A10102251
c 127 10.6 55.8 50 1 A10102252
c 128 10.6 55.8 50 1 A10102252
c 129 10.6 55.8 50 8 A2759877
c 130 10.6 55.8 50 8 CC156374
c 131 10.6 55.8 50 8 CC178643
c 132 10.6 55.8 50 8 CC178647
c 133 10.6 55.8 50 8 CC183595
c 134 10.6 55.8 50 8 CC183642
c 135 10.6 55.8 50 8 CC200501
c 136 10.6 55.8 50 8 CC200515
c 137 10.6 55.8 50 8 CC200522
c 138 10.6 55.8 50 8 CC235392
c 139 10.6 55.8 50 8 CC235393
c 140 10.6 55.8 50 8 CC325402
c 141 10.6 55.8 50 8 CC325432
c 142 10.6 55.8 50 8 CC325540
c 143 10.6 55.8 50 8 CC326063
c 144 10.6 55.8 50 9 CG722613
c 145 10.6 55.8 51 1 A1437786
c 146 10.6 55.8 51 4 BG477735
c 147 10.6 55.8 51 8 CC200483
c 148 10.6 55.8 51 8 CC200494
c 149 10.6 55.8 51 8 CC249684
c 150 10.6 55.8 51 8 CC325545
```

ALIGNMENTS

```
RESULT 1
AA776363/c 55 bp mRNA linear EST 05-FEB-1998
LOCUS ah1ld05.g1 Gessler Wilms tumor Homo sapiens cDNA clone
DEFINITION IMAGE:1156329 3', similar to SW:COX1 HUMAN P00395 CYTOCHROME C
OXIDASE POLYPEPTIDE I ; mRNA sequence.
AA776363
VERSION AA776363.1 GI:2835697
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 55)
AUTHORS Hillier,L., Allen,M., Bowles,L., Dubuque,T., Geisler,G., Jost,S.,
```

TITLE
JOURNAL
COMMENT

Krizman,D., Kucaba,T., Lacy,M., Le,N., Lennon,G., Marra,M.,
Martin,J., Moore,B., Schellenberg,K., Steptoe,M., Tan,F.,
Theising,B., White,Y., Wylie,T., Waterston,R. and Wilson,R.
WashU-NCI human EST Project
Unpublished (1997)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.edu
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Possible reversed clone: similarity on wrong strand
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 1.

FEATURES

source

1..55
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1156329"
/sex="pooled (6)"
/lab_host="DH10B"
/clone_lib="Gessler Wilms tumor"
/note="Vector: pSPORT1; Site 1: SalI; Site 2: NotI; RNA
was prepared from a pool of 6 anonymous Wilms' tumor RNAs.
RNA was prepared by acid-phenol, followed by one round of
oligo dt selection. cDNA library preparation was with
the BRL/life Tech. Superscript Plasmid system. An
oligo-dt NotI primer for first strand synthesis generated
9599cggccc(t)n at the 3' end of the clones. A 5' SalI
adaptor was used with sequence 5'-gtcagccacgcgtccg-3'.
Resulting cDNAs were size selected (average size 2 kb),
NotI digested, and ligated into NotI/SalI-cut pSPORT1.
Library was constructed by Dr. Manfred Gessler."

ORIGIN

Query Match 72.6%; Score 13.8; DB 1; Length 55;
Best Local Similarity 88.2%; Pred. No. 1.5e+04;
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGGTG 17

Db 17 CGGAATGCCCGCACGT 1

RESULT 2

AV836760/c 39 bp mRNA linear EST 09-MAY-2002
LOCUS AV836760 K. Sato unpublished cDNA library: Hordeum vulgare subsp.
DEFINITION vulgare seedling leaves second leaf stage Hordeum vulgare subsp.
vulgare cDNA clone basd2b22, mRNA sequence.

ACCESSION AV836760.1 GI:14528849

VERSION AV836760

KEYWORDS EST.

SOURCE

ORGANISM

Hordeum vulgare subsp. vulgare
Hordeum vulgare subsp. vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Hordeum.

REFERENCE 1 (bases 1 to 39)

Sato,K.

AUTHORS

Barley EST sequencing project in NIG and Okayama Univ

Unpublished (2001)

JOURNAL

COMMENT

Contact: Kazuhiro Sato

Research Institute for Bioreources

Okayama University, Barley Germplasm Center

Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan

Email: kazsato@rib.okayama-u.ac.jp,

URL: http://www.rib.okayama-u.ac.jp/barley/

database: <http://www.shigen.nig.ac.jp/barley/Barley.html>.

```

FEATURES
  source
    Location/Qualifiers
      1..39
        /organism="Hordeum vulgare subsp. vulgare"
        /mol_type="mRNA"
        /cultivar="Haruna Nijo"
        /sub_species="vulgare"
        /db_xref="taxon:112509"
        /clone="basd2b22"
        /tissue_type="seedling leaves"
        /dev_stage="second leaf stage"
        /clone_lib="K. Sato unpublished cDNA library; Hordeum
        vulgare subsp. vulgare seedling leaves second leaf stage"
ORIGIN
  Query Match      70.5%; Score 13.4; DB 1; Length 39;
  Best Local Similarity 87.5%; Pred. No. 2.4e+04;
  Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy  3 GAATGCCCGCGTGT 18
    |||||
Db  30 GAATGNC CGCATGT 15

```

```

RESULT 3
AV841787/c
LOCUS
DEFINITION
  AV841787 Nori Satoh unpublished cDNA library, egg Ciona
  intestinalis cDNA clone rieig06a06, mRNA sequence.
ACCESSION
  AV841787
VERSION
  AV841787.1 GI:16785938
KEYWORDS
  EST.
SOURCE
  Ciona intestinalis
  ORGANISM
    Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
    Phlebobranchia; Clonidae; Ciona.

```

```

REFERENCE
  1 (bases 1 to 54)
  Satoh, N., Satou, Y., Kohara, Y. and Shin-i, T.
  Expressed genes in Ciona intestinalis
  JOURNAL
    Unpublished (2000)
  COMMENT
    Contact: Nori Satoh
    Department of Zoology
    Kyoto University
    Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
    Tel: 81-75-753-4081
    Fax: 81-75-705-1113
    Email: satoheascidian.zool.kyoto-u.ac.jp.

```

```

FEATURES
  source
    Location/Qualifiers
      1..54
        /organism="Ciona intestinalis"
        /mol_type="mRNA"
        /db_xref="taxon:7719"
        /clone="rieig06a06"
        /tissue_type="whole animal"
        /dev_stage="egg"
        /clone_lib="Nori Satoh unpublished cDNA library, egg"
ORIGIN
  Query Match      69.5%; Score 13.2; DB 1; Length 54;
  Best Local Similarity 83.3%; Pred. No. 3.1e+04;
  Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy  1 CGGAATGCCCGCGTGT 18
    |||||
Db  37 CAGATGACCGCGTTT 20

```

```

RESULT 4
BM396904/c
LOCUS
DEFINITION
  BM396904 30 bp mRNA linear EST 17-JAN-2002
  5009-0-26-F02.t.1 Chilcoat/Turkewitz cDNA (large fraction)
  Tetrahymena thermophila cDNA, mRNA sequence.
ACCESSION
  BM396904

```

```

VERSION
  BM396904.1 GI:18196957
KEYWORDS
  EST.
SOURCE
  Tetrahymena thermophila
  ORGANISM
    Tetrahymena thermophila
    Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;
    Hymenostomatida; Tetrahymenina; Tetrahymenidae; Tetrahymena.
  1 (bases 1 to 30)
  Turkewitz, A.P., Karrer, K.M., Jahn, C., Orias, E., Kirk, K.E.,
  Frankel, J. and Klobutcher, L.
  EST from Tetrahymena thermophila, strain CU428.1, growing cells
  JOURNAL
    Unpublished (2002)
  COMMENT
    Contact: Turkewitz AP
    Molecular Genetics and Cell Biology
    University of Chicago
    920 E. 58th Street, Chicago, IL 60637, USA
    Tel: 773 702 4374
    Fax: 773 702 3172
    Email: apturkew@midway.uchicago.edu
    Seq primer: T3.

```

```

FEATURES
  source
    Location/Qualifiers
      1..30
        /organism="Tetrahymena thermophila"
        /mol_type="mRNA"
        /strain="CU428.1"
        /db_xref="taxon:5911"
        /clone_lib="Chilcoat/Turkewitz cDNA (large fraction)"
        /notes="Vector: Bluescript2 SK+; Details on library
        preparation can be found in Chilcoat and Turkewitz (2001)
        Proc. Natl. Acad. Sci USA, 98: 8709-8713."
ORIGIN
  Query Match      68.4%; Score 13; DB 4; Length 30;
  Best Local Similarity 100.0%; Pred. No. 3.9e+04;
  Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  4 AATGCCCCCGCGTG 16
    |||||
Db  29 AATGCCCCCGCGTG 17

```

```

RESULT 5
AV951548/c
LOCUS
DEFINITION
  AV951548 Nori Satoh unpublished cDNA library, egg Ciona
  intestinalis cDNA clone cieg04i05 5', mRNA sequence.
ACCESSION
  AV951548
VERSION
  AV951548.1 GI:19439847
KEYWORDS
  EST.
SOURCE
  Ciona intestinalis
  ORGANISM
    Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
    Phlebobranchia; Clonidae; Ciona.
  1 (bases 1 to 54)
  Satoh, N., Satou, Y., Kohara, Y. and Shin-i, T.
  Expressed genes in Ciona intestinalis
  JOURNAL
    Unpublished (2000)
  COMMENT
    Contact: Nori Satoh
    Department of Zoology
    Kyoto University
    Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
    Tel: 81-75-753-4081
    Fax: 81-75-705-1113
    Email: satoheascidian.zool.kyoto-u.ac.jp.

```

```

FEATURES
  source
    Location/Qualifiers
      1..54
        /organism="Ciona intestinalis"
        /mol_type="mRNA"
        /db_xref="taxon:7719"
        /clone="cieg04i05"
        /tissue_type="whole animal"
        /dev_stage="egg"
        /clone_lib="Nori Satoh unpublished cDNA library, egg"
ORIGIN

```

```

Query Match      67.4%; Score 12.8; DB 2; Length 54;
Best Local Similarity 87.5%; Pred. No. 5.1e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGTG 16
    |||||
Db 19 CGAAATGCCCTCGTG 4

RESULT 6
BG717269
LOCUS
DEFINITION 602689583P1 NIH_MGC_97 Homo sapiens cDNA clone IMAGE:4821885 5',
          mRNA sequence.
ACCESSION BG717269
VERSION BG717269.1 GI:13996456
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 36)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
Toshiyuki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: L1AM10729 row: e column: 22
High quality sequence stop: 36.
Location/Qualifiers
FEATURES
    source
        1..36
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="IMAGE:4821885"
            /lab_host="DH10B"
            /clone_lib="NIH_MGC_97"
            /note="Organ: testis; Vector: pBluescriptR (modified
            pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI
            (gtcgag); Oligo-dT primed using primer
            5'-TTTTTTTTTTTTTTVN-3', size-selected for average
            insert size 2.2 kb and normalized to ROT 5. This is a
            primary library enriched for full-length clones and
            constructed using the Cap-trapper method (Carninci, in
            preparation). Library constructed by M. Brownstein
            (NIMH/NHGRI, National Institutes of Health). Note: this is
            a NIH_MGC Library."

Query Match      66.3%; Score 12.6; DB 4; Length 36;
Best Local Similarity 78.9%; Pred. No. 6.4e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGTG 19
    |||||
Db 3 CGGAGGCCCGCGGCTTG 21

RESULT 7
BG722105
LOCUS
DEFINITION 602689519P1 NIH_MGC_97 Homo sapiens cDNA clone IMAGE:4830340 5',
          mRNA sequence.
ACCESSION BG722105

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```

VERSION BG722105.1 GI:14001292
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 37)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
Toshiyuki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: L1AM10751 row: f column: 05
High quality sequence stop: 37.
Location/Qualifiers
FEATURES
    source
        1..37
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="IMAGE:4830340"
            /lab_host="DH10B"
            /clone_lib="NIH_MGC_97"
            /note="Organ: testis; Vector: pBluescriptR (modified
            pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI
            (gtcgag); Oligo-dT primed using primer
            5'-TTTTTTTTTTTTTTVN-3', size-selected for average
            insert size 2.2 kb and normalized to ROT 5. This is a
            primary library enriched for full-length clones and
            constructed using the Cap-trapper method (Carninci, in
            preparation). Library constructed by M. Brownstein
            (NIMH/NHGRI, National Institutes of Health). Note: this is
            a NIH_MGC Library."

ORIGIN
Query Match      66.3%; Score 12.6; DB 4; Length 37;
Best Local Similarity 78.9%; Pred. No. 6.4e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGTG 19
    |||||
Db 3 CGGAGTGACCCGCGCTTG 21

RESULT 8
A1745090/c
LOCUS
DEFINITION A1745090.x1 NCI_CGAP_OV23 Homo sapiens cDNA clone IMAGE:2218861 3',
          similar to SW:CA12 MOUSE P28481 PROCOLLAGEN ALPHA 1(I) CHAIN
          PRECURSOR [CONTAINS: CHONDROCALCIN]. [3] TR:062031 TR:062032
          ;contains element MSRI repetitive element ;, mRNA sequence.
ACCESSION A1745090
VERSION A1745090.1 GI:5113378
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 40)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

```

Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality

Seq primer: -40UP from Gibco

High quality sequence stop: 1.

Location/Qualifiers

FEATURES

source

1. .40
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2218861"
/tissue_type="tumor, 5 pooled (see description)"
/lab_hosts="DHI08"
/clone_lib="NCI_CGAP_Ov23"
/note="Organ: ovary; Vector: pCMV-SPORT6; Site 1: SalI;
Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt.
Average insert size 1.35 kb. Tumor types include: mixed
Mullerian tumor, papillary serous, clear cell, spindle
cell. All are primary tumors, metastasis positive. Life
Technologies catalog #: 11534-013"

ORIGIN

Query Match 66.3%; Score 12.6; DB 1; Length 40;
Best Local Similarity 78.9%; Pred. No. 6.4e+04;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Cy 1 CGGAATGCCCGCGGTGTTG 19

Db 26 CGGATTCGCCCGGTGTTG 8

RESULT 9

BX892307/c

LOCUS

DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-600H08-023961,
genomic survey sequence.

ACCESSION BX892307

VERSION BX892307.1

KEYWORDS GI:39924802

SOURCE GSS.

ORGANISM Arabidopsis thaliana (thale cress)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE

1

Li, Y., Rosso, M.G., Strizhov, N., Viehoever, P. and Weissshaar, B.

GABI-Kat SimpleSearch: a flanking sequence tag (FST) database for
the identification of T-DNA insertion mutants in Arabidopsis
thaliana

JOURNAL Bioinformatics 19 (11), 1441-1442 (2003)

MEDLINE 22755829

PUBMED 12874060

REFERENCE

2

Rosso, M.G., Li, Y., Strizhov, N., Reiss, B., Dekker, K. and

Weissshaar, B.

An Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
flanking sequence tag-based reverse genetics

JOURNAL Plant Mol. Biol. 53 (1-2), 247-259 (2003)

MEDLINE 23117147

PUBMED 14756321

REFERENCE

3

Strizhov, N., Li, Y., Rosso, M.G., Viehoever, P., Dekker, K.A. and

Weissshaar, B.

High-throughput generation of sequence indexes from T-DNA
mutagenized Arabidopsis thaliana lines

JOURNAL Biotechniques 35 (6), 1164-1168 (2003)

PUBMED 14682050

REFERENCE

4 (bases 1 to 58)

Roos, M.G., Li, Y., Strizhov, N. and Weissshaar, B.

AUTHORS Direct Submission

JOURNAL Submitted (31-MAR-2004)

WEISSHAAR B., Max-Planck-Institut fuer

Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

This sequence has been recovered from the left border of the T-DNA.

It indicates an insertion within the locus defined by BAC clone

T10P12. Details on the protocols used for generation of the

sequence are described in References 1-3. The sequences are

generated at the MPI for Plant Breeding Research in the context of

the GABI-Kat project. GABI-Kat is part of the German Plant Genomics

program designated 'GABI'. Information on line availability can be

found at: <http://www.mpiz-koeln.mpg.de/GABI-kat/>.

Location/Qualifiers

source

1. .58

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="GK-600H08-023961"

/clone_lib="Arabidopsis thaliana T-DNA insertion lines"

/ecotype="Col-0"

/note="PCR was performed on DNA from Arabidopsis thaliana

plants (Ti) which were transformed with the T-DNA from

vector pAC161 (GenBank accession number: AJ537514). The

lines contain one or more T-DNA insertions. The DNA

fragment(s) resulting from the PCR were directly sequenced

to determine the genomic sequence flanking the insertion.

T-DNA derived sequences were removed."

Query Match 66.3%; Score 12.6; DB 9; Length 58;

Best Local Similarity 78.9%; Pred. No. 6.5e+04;

Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Cy 1 CGGAATGCCCGCGGTGTTG 19

Db 40 CGGGAAGTCCCTCGTGTG 22

RESULT 10

BX398127/c

LOCUS

DEFINITION 5009-0-41-A08.t.1 Chilcoat/Turkewitz cDNA (large fraction)-

Tetrahymena thermophila cDNA, mRNA sequence.

ACCESSION BM398127

VERSION BM398127.1

KEYWORDS GI:18198180

SOURCE EST.

ORGANISM Tetrahymena thermophila

Tetrahymena thermophila

Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;

Hymenostomatida; Tetrahymenina; Tetrahymenidae; Tetrahymena.

1 (bases 1 to 30)

Turkewitz, A.P., Karrer, K.M., Jahn, C., Orlas, E., Kirk, K.E.,

Frankel, J. and Klobutcher, L.

EST from Tetrahymena thermophila, strain CU428.1, growing cells

Unpublished (2002)

Contact: Turkewitz AP

Molecular Genetics and Cell Biology

University of Chicago

920 E. 58th Street, Chicago, IL 60637, USA

Tel: 773 702 4374

Fax: 773 702 3172

Email: apturkew@midway.uchicago.edu

Seq primer: T3.

Location/Qualifiers

source

1. .30

/organism="Tetrahymena thermophila"

/mol_type="mRNA"

/strain="CU428.1"

/db_xref="taxon:5911"

/clone_lib="Chilcoat/Turkewitz cDNA (large fraction)"

/note="Vector: Bluescript2 SK+; Details on library

(web address: www.resgen.com) (email contact: info@resgen.com) and
 RessourcenZentrumPrimaarDatenbank, Berlin, Germany (web address:
www.rzpd.de)

Trace considered overall poor quality

Possible reversed clone: similarity on wrong strand

Seq primer: T3 ET from AmerSham

High quality sequence stop: 1.

Location/Qualifiers

FEATURES

source

1..52
 /organism="Danio rerio"
 /mol_type="mRNA"
 /db_xref="taxon:7955"
 /clone="IMAGE:3729257"
 /sex="mixed"
 /tissue_type="26 somite embryos, adult livers, shield
 stage embryos"
 /lab_host="XLI-blue MRF"
 /clone_lib="Zebrafish WashU MPIMG EST"
 /note="Vector: pSPORT1; Site 1: NotI; Site 2: SalI; 1st
 strand cDNA was primed with a Not I - oligo(dT)15 primer
 [5'GCAGTACTTAGTCGCGAGCGCGCCCTTTTCTTTT3'];
 double-stranded cDNA was ligated to Sal I adaptors (BRL),
 digested with Not I and cloned into the Not I and Sal I
 sites of the pSPORT1 vector (BRL). Library was constructed
 by Matthew Clark (Lehrach lab; ICRF, London and Max Planck
 Institut fuer Molekulare Genetik, Berlin). cDNAs for EST
 analysis were selected following oligonucleotide
 hybridization fingerprinting of arrayed clones from
 zebrafish late somitogenesis (26 ss), adult liver or
 embryonic shield stage (5.6 h) libraries. Fingerprint
 data were used to computationally cluster cDNAs, and a
 single cDNA from each cluster was chosen for sequencing.
 In some cases multiple members of the same cluster were
 sequenced to assess clustering parameters or single clones
 were sequenced additional times to assess quality
 control."

ORIGIN

Query Match 62.1%; Score 11.8; DB 1; Length 52;
 Best Local Similarity 86.7%; Pred. No. 1.7e+05;
 Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 4 AATGCCCGCGTGTT 18
 |||||
 Db 38 AATGCCCGCGTGCT 52

RESULT 17
 CR770268
 LOCUS CR770268 55 bp DNA linear GSS 15-SEP-2004
 DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-181E08-013603,
 genomic survey sequence.

ACCESSION CR770268
 VERSION CR770268.1 GI:52138206
 KEYWORDS GSS.
 SOURCE Arabidopsis thaliana (thale cress)
 ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1
 Li, Y., Rosso, M.G., Strizhov, N., Viehoever, P. and Weissshaar, B.
 GABI-Kat SimpleSearch: a flanking sequence tag (EST) database for
 the identification of T-DNA insertion mutants in Arabidopsis
 thaliana

JOURNAL Bioinformatics 19 (11), 1441-1442 (2003)
 MEDLINE 22755829
 PUBMED 12874060

REFERENCE 2
 Rosso, M.G., Li, Y., Strizhov, N., Reiss, B., Dekker, K. and
 Weissshaar, B.

TITLE An Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
 flanking sequence tag-based reverse genetics

JOURNAL

MEDLINE

PUBMED

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Plant Mol. Biol. 53 (1-2), 247-259 (2003)

23117147

14756321

3

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Strizhov, N., Li, Y., Rosso, M.G., Viehoever, P., Dekker, K.A. and

Weissshaar, B.

High-throughput generation of sequence indexes from T-DNA

mutagenized Arabidopsis thaliana lines

BioTechniques 35 (6), 1164-1168 (2003)

23044198

14682050

4

AUTHORS

TITLE

JOURNAL

COMMENT

Li, Y., Strizhov, N., Rosso, M.G. and Weissshaar, B.

Direct Submission

Submitted (15-SEP-2004) Weissshaar, B., Max-Planck-Institut fuer

Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

This sequence has been recovered from the left border of the T-DNA.

It indicates an insertion within the locus defined by BAC clone

T28G19. Details on the protocols used for generation of the

sequence are described in References 1-3. The sequences are

generated at the MPI for Plant Breeding Research in the context of

the GABI-Kat project. GABI-Kat is part of the German Plant Genomics

program designated 'GABI'. Information on line availability can be

found at: <http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

Location/Qualifiers

1..55

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="GK-181E08-013603"

/clone_lib="Arabidopsis thaliana T-DNA insertion lines"

/ecotype="Col-0"

/note="PCR was performed on DNA from Arabidopsis thaliana

plants (T1) which were transformed with the T-DNA from

vector pAC161 (GenBank accession number: AJ537514). The

lines contain one or more T-DNA insertions. The DNA

fragment(s) resulting from the PCR were directly sequenced

to determine the genomic sequence flanking the insertion.

T-DNA derived sequences were removed."

ORIGIN

Query Match 62.1%; Score 11.8; DB 9; Length 55;

Best Local Similarity 86.7%; Pred. No. 1.7e+05;

Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GGAATGCCCGCGGTG 16

|||||

Db 1 GGGATGCCCGCACGTG 15

|||||

RESULT 18

AZ333170/c

LOCUS AZ333170 26 bp DNA linear GSS 29-SEP-2000

DEFINITION 1M062B07F Mouse 10kb plasmid UUGC1M library Mus musculus genomic

clone UUGC1M062B07 F, genomic survey sequence.

ACCESSION AZ333170

VERSION AZ333170.1 GI:10397527

KEYWORDS GSS.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 26)

Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,

Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,

Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von

Niederhausern, A. and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

JOURNAL

MEDLINE

PUBMED

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0062 row: B column: 07
Seq primer: CGTTGTAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 26.

FEATURES

source

Location/Qualifiers
1. .26
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGCIM0062B07"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/clone_lib="Mouse 10kb plasmid UUGCIM library"
/notes="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 [gi|4732114|gb|AF129072.1], a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

Query Match 61.1%; Score 11.6; DB 8; Length 26;
Best Local Similarity 77.8%; Pred. No. 2.1e+05;
Matches 14; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

ORIGIN

Qy 2 GGATGCCCCCGGTGTG 19
|||||
Db 22 GGAAAGCCCCCGTGTGG 5

RESULT 19

LOCUS CO781458 48 bp mRNA linear EST 05-AUG-2004
DEFINITION BL012C.A08 6-Day Axolotl Tail Blastema (6DAXBL) Ambystoma mexicanum cDNA 5' similar to hypothetical protein, mRNA sequence.

ACCESSION CO781458

VERSION CO781458.1 GI:50997438

KEYWORDS EST.

SOURCE Ambystoma mexicanum (axolotl)

ORGANISM Ambystoma mexicanum

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Amphibia; Batrachia; Caudata; Salamandroidea; Ambystomatidae; Ambystoma.

1 (bases 1 to 48)

REFERENCE
AUTHORS Habermann, B., Bebin, A.G., Herklotz, S., Volkmer, M., Eckelt, K., Pehle, K., Epperlein, H.H., Schacker, H.K., Wiese, G. and Tanaka, E.M.
TITLE An Ambystoma mexicanum EST sequencing project: Analysis of 17,352 expressed sequence tags from embryonic and regenerating blastema cDNA libraries

JOURNAL Genome Biol. (2004) In press

COMMENT Contact: Ely M. Tanaka

Tanaka Lab

Max Planck Institute of Molecular Cell Biology and Genetics,
Dresden
Pfotenhauserstrasse 108, 01307 Dresden, Germany
Tel: 0049 351 210 2620
Fax: 0049 351 210 1489
Email: tanaka@mpi-cbg.de
Plate: BL012C row: 08 column: A
Seq primer: GCA CAT TAG GCC TAT TTA GGT GAC A.

FEATURES

source

Location/Qualifiers
1. .48
/organism="Ambystoma mexicanum"
/mol_type="mRNA"
/db_xref="taxon:8296"
/tissue_type="Tail Blastema"
/cell_type="regenerating tail blastema"
/clone_lib="6-Day Axolotl Tail Blastema (6DAXBL)"
/notes="Vector: pCMVSpot6; Site 1: NotI; Site 2: SalI;
Unnormalized cDNA plasmid library prepared by Invitrogen.
Size fractionated mRNA was polydT primed and cloned into NotI-SalI site of pCMVSpot6. Bacterial host is EMDH108-TONA. Average insert size is 1.67 kb.
TAG_Lib=6DAXBL"

ORIGIN

Query Match 61.1%; Score 11.6; DB 7; Length 48;
Best Local Similarity 77.8%; Pred. No. 2.2e+05;
Matches 14; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGGTGT 18

|||||

Db 3 CGGAATGCCCGGTGT 20

RESULT 20

LOCUS BZ662469

DEFINITION SALK_025955.37.60.x Arabidopsis thaliana TDNA insertion lines

Arabidopsis thaliana genomic clone SALK_025955.37.60.x, genomic survey sequence.

ACCESSION BZ662469

VERSION BZ662469.1 GI:28176722

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

1 (bases 1 to 51)

REFERENCE Alonso, J.M., Leisner, T.J., Barajas, P., Chen, H., Cheuk, R.,

Gadrinab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L.,

Shinn, P., Zimmermann, J. and Ecker, J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished (2001)

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGNAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@salk.edu

This is single pass sequence recovered from the left border of

TDNA. This sequence lies within an annotated intron of At3g12520.

Class: TDNA tagged.

FEATURES

source

Location/Qualifiers
1. .51
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Col-0"
/db_xref="taxon:3702"
/clone="SALK_025955.37.60.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines

each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html

ORIGIN

Query Match 61.1%; Score 11.6; DB 8; Length 51;
 Best Local Similarity 77.8%; Pred. No. 2.2e+05;
 Matches 14; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTGT 18
 | ||||| |||||
 Db 6 CTGAATGCCCGTGT 23

RESULT 21
 BM398263/c
 LOCUS 27 bp mRNA linear EST 17-JAN-2002
 DEFINITION 5099-0-43-A08.t.1 Chilcoat/Turkewitz cDNA (large fraction)
 Tetrahymena thermophila cDNA, mRNA sequence.
 ACCESSION BM398263
 VERSION BM398263.1 GI:18198316
 KEYWORDS EST.
 SOURCE Tetrahymena thermophila
 ORGANISM Tetrahymena thermophila
 Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;
 Hymenostomatida; Tetrahymenina; Tetrahymenidae; Tetrahymena.
 REFERENCE 1 (bases 1 to 27)
 AUTHORS Turkewitz,A.P., Karrer,K.M., Jahn,C., Orlas,E., Kirk,K.E.,
 Frankel,J. and Klobutcher,L.
 TITLE EST from Tetrahymena thermophila, strain CU428.1, growing cells
 JOURNAL Unpublished (2002)
 COMMENT Contact: Turkewitz AP
 Molecular Genetics and Cell Biology
 University of Chicago
 920 E. 58th Street, Chicago, IL 60637, USA
 Tel: 773 702 4374
 Fax: 773 702 3172
 Email: apturkew@midway.uchicago.edu
 Seq primer: T3.

FEATURES
source

1..27
 /organism="Tetrahymena thermophila"
 /mol_type="mRNA"
 /strain="CU428.1"
 /db_xref="taxon:5911"
 /clone_libs="Chilcoat/Turkewitz cDNA (large fraction)"
 /note="Vector: Bluescript2 SK+; Details on library preparation can be found in Chilcoat and Turkewitz (2001)
 Proc. Natl. Acad. Sci USA, 98: 8709-8713."

ORIGIN

Query Match 60.0%; Score 11.4; DB 4; Length 27;
 Best Local Similarity 92.3%; Pred. No. 2.7e+05;
 Matches 12; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 AATGCCCGCGTG 16
 | ||||| |||||
 Db 26 AGTGCCCGCGTG 14

RESULT 22
 BM396162/c
 LOCUS 30 bp mRNA linear EST 17-JAN-2002
 DEFINITION 5099-0-18-B05.t.1 Chilcoat/Turkewitz cDNA (large fraction)
 Tetrahymena thermophila cDNA, mRNA sequence.
 ACCESSION BM396162
 VERSION BM396162.1 GI:18196215
 KEYWORDS EST.
 SOURCE Tetrahymena thermophila
 ORGANISM Tetrahymena thermophila
 Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;

Hymenostomatida; Tetrahymenina; Tetrahymenidae; Tetrahymena.
 1 (bases 1 to 30)
 REFERENCE Turkewitz,A.P., Karrer,K.M., Jahn,C., Orlas,E., Kirk,K.E.,
 Frankel,J. and Klobutcher,L.
 TITLE EST from Tetrahymena thermophila, strain CU428.1, growing cells
 JOURNAL Unpublished (2002)
 COMMENT Contact: Turkewitz AP
 Molecular Genetics and Cell Biology
 University of Chicago
 920 E. 58th Street, Chicago, IL 60637, USA
 Tel: 773 702 4374
 Fax: 773 702 3172
 Email: apturkew@midway.uchicago.edu
 Seq primer: T3.

FEATURES
source

1..30
 /organism="Tetrahymena thermophila"
 /mol_type="mRNA"
 /strain="CU428.1"
 /db_xref="taxon:5911"
 /clone_libs="Chilcoat/Turkewitz cDNA (large fraction)"
 /note="Vector: Bluescript2 SK+; Details on library preparation can be found in Chilcoat and Turkewitz (2001)
 Proc. Natl. Acad. Sci USA, 98: 8709-8713."

ORIGIN

Query Match 60.0%; Score 11.4; DB 4; Length 30;
 Best Local Similarity 92.3%; Pred. No. 2.7e+05;
 Matches 12; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 AATGCCCGCGTG 16
 | ||||| |||||
 Db 29 AGTGCCCGCGTG 17

RESULT 23
 BM400993/c
 LOCUS 34 bp mRNA linear EST 17-JAN-2002
 DEFINITION 5099-0-81-B09.t.1 Chilcoat/Turkewitz cDNA (large fraction)
 Tetrahymena thermophila cDNA, mRNA sequence.
 ACCESSION BM400993
 VERSION BM400993.1 GI:18201046
 KEYWORDS EST.
 SOURCE Tetrahymena thermophila
 ORGANISM Tetrahymena thermophila
 Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;
 Hymenostomatida; Tetrahymenina; Tetrahymenidae; Tetrahymena.
 REFERENCE 1 (bases 1 to 34)
 AUTHORS Turkewitz,A.P., Karrer,K.M., Jahn,C., Orlas,E., Kirk,K.E.,
 Frankel,J. and Klobutcher,L.
 TITLE EST from Tetrahymena thermophila, strain CU428.1, growing cells
 JOURNAL Unpublished (2002)
 COMMENT Contact: Turkewitz AP
 Molecular Genetics and Cell Biology
 University of Chicago
 920 E. 58th Street, Chicago, IL 60637, USA
 Tel: 773 702 4374
 Fax: 773 702 3172
 Email: apturkew@midway.uchicago.edu
 Seq primer: T3.

FEATURES
source

1..34
 /organism="Tetrahymena thermophila"
 /mol_type="mRNA"
 /strain="CU428.1"
 /db_xref="taxon:5911"
 /clone_libs="Chilcoat/Turkewitz cDNA (large fraction)"
 /note="Vector: Bluescript2 SK+; Details on library preparation can be found in Chilcoat and Turkewitz (2001)
 Proc. Natl. Acad. Sci USA, 98: 8709-8713."

ORIGIN

Query Match 60.0%; Score 11.4; DB 4; Length 34;

```

Best Local Similarity 85.7%; Pred. No. 2.7e+05;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 GAATGCCCGCGGTG 16
   ||| ||| ||| |||
Db 22 GACTGCGCGCGGTG 9

RESULT 24
AUI05629
LOCUS AUI05629 50 bp mRNA linear EST 28-JAN-2004
DEFINITION AUI05629 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
            HSI02111, mRNA sequence.
ACCESSION AUI05629
VERSION AUI05629.1 GI:13555150
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 50)
AUTHORS Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J.,
            Hata,H., Ota,T., Isogai,T., Tanaka,T., Morishita,S., Okubo,K.,
            Sakaki,Y., Nakamura,Y., Suyama,A. and Sugano,S.
TITLE Diverse transcriptional initiation revealed by fine, large-scale
JOURNAL mapping of mRNA start sites
MEDLINE ENBO Rep. 2 (5), 388-393 (2001)
PUBMED 21270072
COMMENT Contact: Yutaka Suzuki
            Department of Virology
            Institute of Medical Science, University of Tokyo
            4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
            Email: yuzuki@ims.u-tokyo.ac.jp
Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and
Sugano,S. Construction and characterization of a full
length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2),
149-156 (1997).
FEATURES
    source
        1..50
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="HSI02111"
            /clone_lib="Sugano Homo sapiens cDNA library"
ORIGIN
Query Match 60.0%; Score 11.4; DB 1; Length 50;
Best Local Similarity 92.3%; Pred. No. 2.8e+05;
Matches 12; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 TGCCCGCGCGTGT 18
   ||| ||| ||| |||
Db 25 TGCCCGCGCGTGT 37

RESULT 25
A2769819
LOCUS A2769819 27 bp DNA linear GSS 16-FEB-2001
DEFINITION IM0570E20R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
            clone UUGC1M0570E20 R, genomic survey sequence.
ACCESSION A2769819
VERSION A2769819.1 GI:12890359
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
            1 (bases 1 to 27)
REFERENCE 1
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
            Irlam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T.,
            Rilly,M., Rose,R., Rose,R., Stokes,R., Tingey,A., von
            Niederhausern,A. and Wright,D.,Weiss,R.

```

```

TITLE Mouse whole genome scaffolding with paired end reads from 10kb
JOURNAL plasmid inserts
COMMENT Unpublished (2000)
        Contact: Robert B. Weiss
        University of Utah Genome Center
        University of Utah
        Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
        84112, USA
        Tel: 801 585 5606
        Fax: 801 585 7177
        Email: ddunn@genetics.utah.edu
        Insert Length: 10000 Std Error: 0.00
        Plate: 0570 row: E column: 20
        Seq primer: CACACAGAAACAGCTATGACC
        Class: plasmid ends
        High quality sequence stop: 27.
FEATURES
    Location/Qualifiers
        1..27
            /organism="Mus musculus"
            /mol_type="genomic DNA"
            /strain="C57BL/6J"
            /db_xref="taxon:10090"
            /clone="UUGC1M0570E20"
            /sex="Male"
            /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
            /notes="Vector: PWD42nv; Purified genomic DNA from M.
            musculus C57BL/6J (male) was obtained from the Jackson
            Laboratory Mouse DNA Resource
            (http://www.jax.org/resources/documents/dnares/). The DNA
            was hydrodynamically sheared by repeated passage through a
            0.005 inch orifice at constant velocity. The sheared DNA
            was blunt end-repaired with T4 DNA polymerase and T4
            polynucleotide kinase. Adaptor oligonucleotides were
            ligated to the blunt ends in high molar excess. The
            adaptor DNA was purified and size-selected for a 9.5 to
            10.5 kb range using preparative agarose gel
            electrophoresis. Vector DNA was prepared from a derivative
            of pWD42 (GI4732114|GB|AF129072.1), a copy-number
            inducible derivative of plasmid R1. The vector was ligated
            with adaptors complementary to the insert adaptors and
            purified. The sheared, adaptor mouse DNA was annealed to
            adaptor vector DNA, and transformed into
            chemically-competent E. coli XL10-Gold (Stratagene) cells
            and selected for ampicillin resistance."
ORIGIN
Query Match 58.9%; Score 11.2; DB 8; Length 27;
Best Local Similarity 81.2%; Pred. No. 3.5e+05;
Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 AATGCCCGCGGTGTG 19
   ||| ||| ||| |||
Db 11 AATGCCCGCGGTATG 26

RESULT 26
A1001661
LOCUS A1001661 31 bp mRNA linear EST 13-OCT-1999
DEFINITION EST0243 Tilapia brain cDNA library in pUC18 Orochromis niloticus
            cDNA clone 1109, mRNA sequence.
ACCESSION A1001661
VERSION A1001661.1 GI:3201423
KEYWORDS EST.
SOURCE Orochromis niloticus (Nile tilapia)
ORGANISM Orochromis niloticus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
            Acanthomorpha; Acanthopterygii; Percomorpha; Perciformes;
            Labroidae; Cichlidae; Orochromis.
            1 (bases 1 to 31)
REFERENCE 1
AUTHORS Hamilton,L.C., MacPherson,G. and Wright,J.M.
TITLE Expressed sequence tags from a tilapia (Oreochromis niloticus)

```

JOURNAL
COMMENT

brain cDNA library
Unpublished (1998)
Other ESTs: EST0244
Contact: Wright JM
Department of Biology
Dalhousie University
Halifax, NS Canada B3H 4J1
Tel: 902 494 6468
Fax: 902 494 3736
Email: jmwright@dal.ca
Insert Length: 1000 Std Error: 50.00
Seq primer: M13 forward sequencing primer.

FEATURES
source

1. .31
/organism="Oreochromis niloticus"
/mol_type="mRNA"
/db_xref="taxon:8128"
/clone="11109"
/sex="male"
/dev_stage="adult"
/clone_lib="Tilapia brain cDNA library in pUC18"
/note="Organ: brain; Vector: pUC18; cDNA inserts were blunt end cloned into SmaI site of pUC18."

ORIGIN

Query Match 58.9%; Score 11.2; DB 1; Length 31;
Best Local Similarity 81.2%; Pred. No. 3.5e+05;
Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTG 16
|||||
Db 8 CGGAATGCCCGCGG 23

RESULT 27

BM400565/c
LOCUS
DEFINITION
5009-0-75-F09.t.1 Chilcoat/Turkewitz cDNA (large fraction)
Tetrahymena thermophila cDNA, mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE

ORGANISM

REFERENCE
AUTHORS
Turkewitz, A.P., Karrer, K.M., Jahn, C., Orlas, E., Kirk, K.E.,
Frankel, J. and Klobutcher, L.

TITLE
JOURNAL
COMMENT
EST from Tetrahymena thermophila, strain CU428.1, growing cells
Unpublished (2002)
Contact: Turkewitz AP
Molecular Genetics and Cell Biology
University of Chicago
920 E. 58th Street, Chicago, IL 60637, USA
Tel: 773 702 4374
Fax: 773 702 3172
Email: apturkew@midway.uchicago.edu

FEATURES
source

1. .31
/organism="Tetrahymena thermophila"
/mol_type="mRNA"
/strain="CU428.1"
/db_xref="taxon:5911"
/clone_lib="Chilcoat/Turkewitz cDNA (large fraction)"
/note="Vector: Bluescript2 SK+; Details on library preparation can be found in Chilcoat and Turkewitz (2001).
Proc. Natl. Acad. Sci USA, 98: 8709-8713."

ORIGIN

Query Match 58.9%; Score 11.2; DB 4; Length 31;
Best Local Similarity 81.2%; Pred. No. 3.5e+05;

Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
Qy 1 CGGAATGCCCGCGTG 16
|||||
Db 26 CCGTACGCCCGCGTG 11

RESULT 28

BM398283/c
LOCUS
DEFINITION
5009-0-43-D08.t.1 Chilcoat/Turkewitz cDNA (large fraction)
Tetrahymena thermophila cDNA, mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BM398283 33 bp mRNA linear EST 17-JAN-2002
5009-0-43-D08.t.1 Chilcoat/Turkewitz cDNA (large fraction)
Tetrahymena thermophila cDNA, mRNA sequence.
BM398283
BM398283.1 GI:18198336
EST.
Tetrahymena thermophila
Tetrahymena thermophila
Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;
Hymenostomatida; Tetrahymenina; Tetrahymenidae; Tetrahymena.
1 (bases 1 to 33)
Turkewitz, A.P., Karrer, K.M., Jahn, C., Orlas, E., Kirk, K.E.,
Frankel, J. and Klobutcher, L.

EST from Tetrahymena thermophila, strain CU428.1, growing cells
Unpublished (2002)
Contact: Turkewitz AP
Molecular Genetics and Cell Biology
University of Chicago
920 E. 58th Street, Chicago, IL 60637, USA
Tel: 773 702 4374
Fax: 773 702 3172
Email: apturkew@midway.uchicago.edu

Seq primer: T3.

Location/Qualifiers

1. .33

/organism="Tetrahymena thermophila"

/mol_type="mRNA"

/strain="CU428.1"

/db_xref="taxon:5911"

/clone_lib="Chilcoat/Turkewitz cDNA (large fraction)"

/note="Vector: Bluescript2 SK+; Details on library preparation can be found in Chilcoat and Turkewitz (2001).
Proc. Natl. Acad. Sci USA, 98: 8709-8713."

ORIGIN

Query Match 58.9%; Score 11.2; DB 4; Length 33;
Best Local Similarity 81.2%; Pred. No. 3.5e+05;
Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 CGGAATGCCCGCGTG 16
|||||
Db 27 CGGCCAGCCCGCGTG 12

RESULT 29

AU009955/c

LOCUS

DEFINITION

39 bp mRNA linear EST 31-JUL-1998

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Unpublished (1998)

Contact: Mitsunori Morimyo

Genome Research Group

National Institute of Radiological Sciences
9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan
Email: morimyo@nirs.go.jp.
Location/Qualifiers

1. .39
/organism="Schizosaccharomyces pombe"
/mol_type="mRNA"
/strain="972"
/db_xref="taxon:4896"
/clone="spc00715"
/sex="h minus"
/clone_lib="Schizosaccharomyces pombe late log phase cDNA"
/notes="Vector: M13mp19; The cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the SmaI site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, http://www.nirs.go.jp)"

ORIGIN

Query Match 58.9%; Score 11.2; DB 1; Length 39;
Best Local Similarity 81.2%; Pred. No. 3.5e+05;
Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CGGAATGCCCGCGGTG 16
|||||
DB 23 CGGATGCCACGCGCG 8

RESULT 30
AI544749/c
LOCUS
DEFINITION
fb66g01.x1 Zebrafish WashU MPIMG EST Danio rerio cDNA clone
IMAGE:3716880 3' similar to TR:Q13896 Q13896 ALPHA-1 TYPE I
COLLAGEN PRECURSOR ;, mRNA sequence.
AI544749
AI544749.1 GI:4462122
EST.
Danio rerio (zebrafish)
ORGANISM
Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
1 (bases 1 to 46)
Clark, M., Johnson, S.L., Lehrach, H., Lee, R., Li, F., Marra, M.,
Eddy, S., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y.,
Person, B., Swaller, F., Gibbons, M., Pape, D., Harvey, N., Schurk, R.,
Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R.,
Waterston, R. and Wilson, R.
WashU Zebrafish EST Project 1998
Unpublished (1998)
Contact: Stephen L. Johnson
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: zbrfish@watson.wustl.edu
cDNA Library Preparation: Matthew Clark. cDNA Library Arrayed by:
Matthew Clark. DNA Sequencing by: Washington University Genome
Sequencing Center Clone Distribution: Genome Systems, St. Louis,
Missouri (web address: www.genomesystems.com) (email contact:
info@genomesystems.com) and Research Genetics, Huntsville, Alabama
(web address: www.resgen.com) (email contact: info@resgen.com) and
ResourcenZentrumPrimarDatenbank, Berlin, Germany (web address:
www.rzpd.de)
Trace considered overall poor quality
Possible reversed clone: similarity on wrong strand
Seq primer: T7 ET from Amersham
High quality sequence stop: 1
POLYA=No.

FEATURES
source
1. .46
Location/Qualifiers

/organism="Danio rerio"
/mol_type="mRNA"
/db_xref="taxon:7955"
/clone="IMAGE:3716880"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield
stage embryos"
/lab_host="XLI-blue MRF"
/clone_lib="Zebrafish WashU MPIMG EST"
/notes="Vector: pSPORT1; Site 1: NotI; Site 2: SalI; 1st
strand cDNA was primed with a Not I - oligo(dT)15 primer
[5' pGACTAGTCTAGATCGGCGCGCCCTTTTCTTTT3'];
double-stranded cDNA was ligated to Sal I adaptors (BRL),
digested with Not I and cloned into the Not I and Sal I
sites of the pSPORT1 vector (BRL). Library was constructed
by Matthew Clark (Lehrach lab; ICRF, London and Max Planck
Institut fuer Molekulare Genetik, Berlin). cDNAs for EST
analysis were selected following oligonucleotide
hybridization fingerprinting of arrayed clones from
zebrafish late somitogenesis (26 ss), adult liver or
embryonic shield stage (5.6 h) libraries. Fingerprint
data were used to computationally cluster cDNAs, and a
single cDNA from each cluster was chosen for sequencing.
In some cases multiple members of the same cluster were
sequenced to assess clustering parameters or single clones
were sequenced additional times to assess quality
control."

ORIGIN

Query Match 58.9%; Score 11.2; DB 1; Length 46;
Best Local Similarity 81.2%; Pred. No. 3.6e+05;
Matches 13; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GGAATGCCCGCGGTG 17
|||||
DB 34 GGTATGCCCGCGGTG 19

Search completed: September 9, 2005, 22:12:48
Job time : 1783 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 198.688 Seconds
(without alignments)
655.473 Million cell updates/sec

Title: US-10-729-421-35

Perfect score: 22

Sequence: 1 agcccttcagtcacatcaag 22

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database : N_Geneseq_16Dec04.*

1: Geneseq1980s.*

2: Geneseq1990s.*

3: Geneseq2000s.*

4: Geneseq2001as.*

5: Geneseq2001bs.*

6: Geneseq2002as.*

7: Geneseq2002bs.*

8: Geneseq2003as.*

9: Geneseq2003bs.*

10: Geneseq2003cs.*

11: Geneseq2003ds.*

12: Geneseq2004as.*

13: Geneseq2004bs.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|-------------|---------------------|
| 1 | 22 | 100.0 | 22 | ADQ30665 | Adq30665 West Nile |
| 2 | 22 | 100.0 | 24 | ADN36704 | Adn36704 West Nile |
| 3 | 22 | 100.0 | 24 | ADN36702 | Adn36702 West Nile |
| 4 | 22 | 100.0 | 24 | ADN36703 | Adn36703 West Nile |
| 5 | 22 | 100.0 | 51 | ADN36716 | Adn36716 West Nile |
| 6 | 22 | 100.0 | 51 | ADN36714 | Adn36714 West Nile |
| 7 | 22 | 100.0 | 51 | ADN36715 | Adn36715 West Nile |
| 8 | 22 | 100.0 | 69 | ADN36694 | Adn36694 West Nile |
| 9 | 22 | 100.0 | 365 | ABK51710 | Abk51710 Partial c |
| 10 | 22 | 100.0 | 366 | ABQ76684 | Abq76684 WNV Cwt DN |
| 11 | 22 | 100.0 | 967 | ADQ30647 | Adq30647 West Nile |
| 12 | 22 | 100.0 | 10945 | ADR32078 | Adr32078 Genomic D |
| 13 | 22 | 100.0 | 10945 | ADR67768 | Adr67768 West Nile |
| 14 | 22 | 100.0 | 10975 | ADN98022 | Adn98022 West Nile |
| 15 | 22 | 100.0 | 11029 | ABZ68481 | Abz68481 Nucleotid |
| 16 | 22 | 100.0 | 11029 | 10 ABV74821 | Abv74821 West Nile |
| 17 | 22 | 100.0 | 11029 | 12 ADN98023 | Adn98023 West Nile |
| 18 | 21 | 95.5 | 22 | ADN36705 | Adn36705 West Nile |
| 19 | 21 | 95.5 | 49 | ADN36717 | Adn36717 West Nile |
| 20 | 20.4 | 92.7 | 51 | ADN36713 | Adn36713 West Nile |

| | | | | | | |
|----|------|------|--------|----|-------------|--------------------|
| 21 | 20 | 90.9 | 24 | 12 | ADN36706 | Adn36706 West Nile |
| 22 | 20 | 90.9 | 51 | 12 | ADN36718 | Adn36718 West Nile |
| 23 | 19 | 86.4 | 24 | 12 | ADN36701 | Adn36701 West Nile |
| 24 | 18.8 | 85.5 | 10962 | 12 | ADK13681 | Adk13681 West Nile |
| 25 | 18.8 | 85.5 | 44920 | 12 | ADQ97910 | Adq97910 Human can |
| 26 | 18 | 81.8 | 532 | 1 | AAK90221 | Aak90221 Malaria-s |
| 27 | 17.8 | 80.9 | 247 | 3 | AAK09236 | Aak09236 Human sec |
| 28 | 17.8 | 80.9 | 348 | 3 | AAA43855 | Aaa43855 Human sec |
| 29 | 17.8 | 80.9 | 425 | 10 | ADF80038 | Adf80038 Leukaemia |
| 30 | 17.8 | 80.9 | 447 | 2 | AAV49571 | Aav49571 Human lym |
| 31 | 17.8 | 80.9 | 616 | 13 | ADS19174 | Ads19174 Human C-t |
| 32 | 17.8 | 80.9 | 655 | 2 | AAV83108 | Aav83108 Human C-t |
| 33 | 17.8 | 80.9 | 697 | 2 | AAV49570 | Aav49570 Human lym |
| 34 | 17.8 | 80.9 | 697 | 2 | AAV54641 | Aav54641 Nucleotid |
| 35 | 17.8 | 80.9 | 697 | 6 | ABL41987 | AbL41987 Nucleotid |
| 36 | 17.8 | 80.9 | 701 | 10 | ADC38673 | Adc38673 Human CDN |
| 37 | 17.8 | 80.9 | 759 | 6 | ABK84720 | Abk84720 Human CDN |
| 38 | 17.8 | 80.9 | 759 | 6 | ABK48103 | Abk48103 Human CDN |
| 39 | 17.8 | 80.9 | 759 | 10 | ADD18707 | Add18707 Human dis |
| 40 | 17.8 | 80.9 | 759 | 13 | ADR06491 | Adr06491 Human AIC |
| 41 | 17.8 | 80.9 | 759 | 13 | ADP54852 | Adp54852 Human PRO |
| 42 | 17.8 | 80.9 | 935 | 12 | ADQ18422 | Adq18422 Human sof |
| 43 | 17.8 | 80.9 | 1404 | 12 | ADQ22924 | Adq22924 Human sof |
| 44 | 17.8 | 80.9 | 1620 | 13 | ADT46681 | Adt46681 Bacterial |
| 45 | 17.8 | 80.9 | 9375 | 4 | AAK84948 | Aak84948 Human imm |
| 46 | 17.8 | 80.9 | 10301 | 4 | AAK84949 | Aak84949 Human imm |
| 47 | 17.8 | 80.9 | 110000 | 2 | AAV21209_14 | AAV21209_14 |
| 48 | 17.2 | 78.2 | 1420 | 6 | ABT08161 | Abt08161 XisA reco |
| 49 | 17.2 | 78.2 | 1441 | 6 | ABT08163 | Abt08163 NLS-XisA |
| 50 | 17.2 | 78.2 | 1855 | 2 | AAV27359 | Aav27359 Streptoco |
| 51 | 17.2 | 78.2 | 1855 | 6 | ABQ84827 | Abq84827 S. pneumo |
| 52 | 17.2 | 78.2 | 1855 | 10 | ADC45152 | Adc45152 S. pneumo |
| 53 | 17.2 | 78.2 | 3119 | 5 | AAK72586 | Aak72586 DNA encod |
| 54 | 17.2 | 78.2 | 3789 | 13 | ADR93829 | Adr93829 Novel S. |
| 55 | 17.2 | 78.2 | 3840 | 10 | ABX05894 | Abx05894 S. pneumo |
| 56 | 17.2 | 78.2 | 3840 | 12 | ADM91840 | Adm91840 S. pneumo |
| 57 | 17.2 | 78.2 | 4841 | 4 | AAK52955 | Aak52955 Human pol |
| 58 | 17.2 | 78.2 | 4880 | 4 | AAK51971 | Aak51971 Human pol |
| 59 | 17.2 | 78.2 | 4898 | 10 | ABZ79896 | Abz79896 Human nuc |
| 60 | 17.2 | 78.2 | 4915 | 12 | ADQ18215 | Adq18215 Human sof |
| 61 | 17.2 | 78.2 | 4915 | 13 | ADR26068 | Adr26068 Breast ca |
| 62 | 17.2 | 78.2 | 5037 | 12 | ADQ22765 | Adq22765 Human sof |
| 63 | 17.2 | 78.2 | 5309 | 6 | ABT08172 | Abt08172 Recombina |
| 64 | 17.2 | 78.2 | 16080 | 6 | AAD28651 | Aad28651 Human Sal |
| 65 | 17.2 | 78.2 | 16535 | 2 | AAV52207 | AAV52207 Streptoco |
| 66 | 17.2 | 78.2 | 110000 | 10 | ABS56454_01 | ABS56454_01 |
| 67 | 17.2 | 78.2 | 163701 | 13 | ABD33351 | Abd33351 Murine ca |
| 68 | 17 | 77.3 | 17 | 6 | ACN05481 | Acn05481 WNV Amber |
| 69 | 17 | 77.3 | 17 | 6 | ACN09471 | Acn09471 WNV minus |
| 70 | 17 | 77.3 | 17 | 6 | ACN04728 | Acn04728 WNV DNaz |
| 71 | 17 | 77.3 | 17 | 6 | ACN09470 | Acn09470 WNV minus |
| 72 | 17 | 77.3 | 17 | 6 | ACN13599 | Acn13599 WNV minus |
| 73 | 17 | 77.3 | 17 | 6 | ACN12229 | Acn12229 WNV minus |
| 74 | 17 | 77.3 | 17 | 6 | ACN09469 | Acn09469 WNV minus |
| 75 | 17 | 77.3 | 17 | 6 | ACN01443 | Acn01443 WNV inoz |
| 76 | 17 | 77.3 | 17 | 6 | ACN05482 | Acn05482 WNV Amber |
| 77 | 16.8 | 76.4 | 342 | 6 | ABX99067 | Abx99067 Rice endo |
| 78 | 16.8 | 76.4 | 440 | 6 | ABQ55371 | Abq55371 Human ova |
| 79 | 16.8 | 76.4 | 594 | 12 | ACH75379 | Ach75379 Human gen |
| 80 | 16.8 | 76.4 | 742 | 2 | AAV00437 | Aav00437 Clone H90 |
| 81 | 16.8 | 76.4 | 943 | 5 | AAK66880 | Aak66880 DNA encod |
| 82 | 16.8 | 76.4 | 4547 | 6 | AAD28652 | Aad28652 Mouse Sal |
| 83 | 16.8 | 76.4 | 143412 | 11 | ACN44512 | Acn44512 Mouse gen |
| 84 | 16.4 | 74.5 | 518 | 13 | ADQ79152 | Adq79152 Novel can |
| 85 | 16.4 | 74.5 | 650 | 12 | ADK34072 | Adk34072 Yeast let |
| 86 | 16.4 | 74.5 | 656 | 13 | ADQ50257 | Adq50257 Novel can |
| 87 | 16.4 | 74.5 | 960 | 10 | ABX06892 | Abx06892 S. pneumo |
| 88 | 16.4 | 74.5 | 963 | 3 | AAZ46474 | Aaz46474 S. pneumo |
| 89 | 16.4 | 74.5 | 963 | 4 | AAK55886 | Aak55886 Streptoco |
| 90 | 16.4 | 74.5 | 963 | 4 | AAK55543 | Aak55543 Streptoco |
| 91 | 16.4 | 74.5 | 963 | 8 | ACA9937 | AcA9937 Prokaryot |
| 92 | 16.4 | 74.5 | 963 | 13 | ADR91898 | Adr91898 Novel S. |
| 93 | 16.4 | 74.5 | 2643 | 6 | ABN79826 | Abn79826 Fungal ZB |

c 94 16.4 74.5 2646 2 AAG61607 Mutated G
c 95 16.4 74.5 2646 13 ADT47700 Bacterial
c 96 16.4 74.5 2811 12 ADJ92822 Saccharom
c 97 16.4 74.5 3189 2 AAV65242 DNA encod
c 98 16.4 74.5 3694 6 ABK86400 Yeast GAL
c 99 16.4 74.5 3694 12 ADN60220 S. cerevi
c 100 16.4 74.5 3902 2 AAV52345 Streptoco

ALIGNMENTS

RESULT 1

ADQ30665

ID ADQ30665 standard; DNA; 22 BP.

XX

AC ADQ30665;

XX

DT 23-SEP-2004 (first entry)

XX

DE West Nile Virus capsid gene antisense primer WNVVA2.

XX

KW ss; primer; West Nile Virus; diagnosis.

XX

OS West Nile virus.

XX

PN WO2004055159-A2.

XX

XX WO2004055159-A2.

XX

PD 01-JUL-2004.

XX

PF 05-DEC-2003; 2003WO-US038750.

XX

PR 12-DEC-2002; 2002US-0432850P.

XX

PR 20-JUN-2003; 2003US-0480431P.

XX

XX (CHIR) CHIRON CORP.

XX

PI Shyamala V;

XX

DR WPI; 2004-488058/46.

XX

XX New isolated oligonucleotides for accurately diagnosing West Nile virus

XX

PT infection or for capturing, detecting and quantitating West Nile virus in

XX

PT blood samples.

XX

XX Claim 1; SEQ ID NO 35; 56pp; English.

XX

CC The invention relates to an isolated oligonucleotide not more than 60
CC nucleotides in length comprising a nucleotide sequence (S1) of at least
CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
CC 20, 21 or 23 bp) given in the specification derived from the West Nile
CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
CC identity to the nucleotide sequence of (S1), or complements of (S1) and
CC end and/or the 3'-end. The detectable label is a fluorescent label
CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
CC (TAMRA), and 2',4',5',7'-tetrachloro-4,7-dichlorofluorescein (TET). The
CC composition and methods are useful for accurately diagnosing West Nile
CC virus infection or for capturing, detecting and quantitating West Nile
CC virus in biological samples, particularly blood samples. This sequence
CC corresponds to a PCR primer to amplify a fragment of the capsid gene of
CC the WNV genome. The fragment is detected using the oligonucleotides of
CC the invention.

XX

SQ Sequence 22 BP; 6 A; 8 C; 3 G; 5 T; 0 U; 0 Other;

XX

Query Match 100.0%; Score 22; DB 12; Length 22;

XX

Best Local Similarity 100.0%; Pred. No. 0.8; Mismatches 0; Indels 0; Gaps 0;

XX

Matches 22; Conservative 0;

XX

Oy 1 AGCCCTCTTCAGTCCAATCAAG 22

Db 1 AGCCCTCTTCAGTCCAATCAAG 22

RESULT 2

ADN36704

ID ADN36704 standard; DNA; 24 BP.

XX

AC ADN36704;

XX

DT 15-JUL-2004 (first entry)

XX

DE West Nile virus detection-related oligonucleotide probe SeqID26.

XX

KW hybridisation assay probe; nucleic acid detection;

XX

KW target-complementary sequence; flavivirus; West Nile virus; WNV;

XX

KW RNA virus; infection; meningitis; encephalitis;

XX

KW high throughput screening; probe; ss.

XX

OS West Nile virus.

XX

PN WO2004036190-A2.

XX

PD 29-APR-2004.

XX

PF 10-OCT-2003; 2003WO-US033639.

XX

PR 16-OCT-2002; 2002US-0418891P.

XX

PR 25-NOV-2002; 2002US-0429006P.

XX

PR 24-FEB-2003; 2003US-0449810P.

XX

XX (GENP-) GEN-PROBE INC.

XX

PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX

DR WPI; 2004-389590/36.

XX

XX New hybridization assay probe comprising target-complementary sequence of
XX bases, useful in detecting flavivirus, e.g. West Nile virus.

XX

PS Claim 78; SEQ ID NO 26; 135pp; English.

XX

CC This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.

XX

SQ Sequence 24 BP; 7 A; 8 C; 3 G; 6 T; 0 U; 0 Other;

XX

Query Match 100.0%; Score 22; DB 12; Length 24;

XX

Best Local Similarity 100.0%; Pred. No. 0.81; Mismatches 0; Indels 0; Gaps 0;

XX

Matches 22; Conservative 0;

XX

Oy 1 AGCCCTCTTCAGTCCAATCAAG 22

Db 3 AGCCCTCTTCAGTCCAATCAAG 24

XX

RESULT 3

ADN36702

ID ADN36702 standard; DNA; 24 BP.

XX

AC ADN36702;

XX

DT 15-JUL-2004 (first entry)

XX

DE West Nile virus detection-related oligonucleotide probe SeqID24.
 XX hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.
 XX West Nile virus.
 OS
 XX WO2004036190-A2.
 PN
 XX 29-APR-2004.
 PD
 XX 10-OCT-2003; 2003WO-US033639.
 XX
 XX 16-OCT-2002; 2002US-0418891P.
 PR
 XX 25-NOV-2002; 2002US-0429006P.
 PR
 XX 24-FEB-2003; 2003US-0449810P.
 PR
 XX (GENP-) GEN-PROBE INC.
 PA
 XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX WPI; 2004-389590/36.
 DR
 XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 PT
 XX Claim 78; SEQ ID NO 24; 135pp; English.
 PS
 XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.
 XX
 SQ Sequence 24 BP; 7 A; 8 C; 4 G; 5 T; 0 U; 0 Other;
 Query Match 100.0%; Score 22; DB 12; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.81;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 AGCCCTCTTCAGTCCCAATCAAG 22
 DB 1 AGCCCTCTTCAGTCCCAATCAAG 22
 RESULT 4
 ID ADN36703
 ADN36703 standard; DNA; 24 BP.
 XX
 AC ADN36703;
 XX
 XX 15-JUL-2004 (first entry)
 DT
 XX West Nile virus detection-related oligonucleotide probe SeqID25.
 DE
 XX hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.
 XX West Nile virus.
 OS
 XX WO2004036190-A2.
 PN

XX 29-APR-2004.
 PD
 XX 10-OCT-2003; 2003WO-US033639.
 PF
 XX 16-OCT-2002; 2002US-0418891P.
 PR
 XX 25-NOV-2002; 2002US-0429006P.
 PR
 XX 24-FEB-2003; 2003US-0449810P.
 PR
 XX (GENP-) GEN-PROBE INC.
 PA
 XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX WPI; 2004-389590/36.
 DR
 XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 PT
 XX Claim 78; SEQ ID NO 25; 135pp; English.
 PS
 XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.
 XX
 SQ Sequence 24 BP; 6 A; 8 C; 4 G; 6 T; 0 U; 0 Other;
 Query Match 100.0%; Score 22; DB 12; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.81;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 AGCCCTCTTCAGTCCCAATCAAG 22
 DB 2 AGCCCTCTTCAGTCCCAATCAAG 23
 RESULT 5
 ADN36716
 ID ADN36716 standard; DNA; 51 BP.
 XX
 AC ADN36716;
 XX
 XX 15-JUL-2004 (first entry)
 DT
 XX West Nile virus detection-related oligonucleotide probe SeqID38.
 DE
 XX hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.
 XX West Nile virus.
 OS
 XX Enterobacteria phage T7.
 XX
 XX Key Location/Qualifiers
 FT misc_feature 1..27
 FT /tag= a
 FT /note= "T7 promoter sequence"
 FT misc_feature 28..51
 FT /tag= b
 FT /note= "WNV-complementary sequence"
 XX
 XX WO2004036190-A2.
 PN
 XX

```
PD 29-APR-2004.
XX
XX 10-OCT-2003; 2003WO-US033639.
XX
XX 16-OCT-2002; 2002US-0418891P.
XX
XX 25-NOV-2002; 2002US-0429006P.
XX
XX 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
XX
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
XX bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Disclosure; SEQ ID NO 38; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
XX detecting a nucleic acid, which is a probe sequence that comprises a
XX target-complementary sequence of bases, and optionally one or more base
XX sequences that are not complementary to the nucleic acid that is to be
XX detected. The hybridisation assay probes and the kits are useful in
XX detecting and amplifying a target nucleic acid sequence, for example
XX flavivirus like West Nile virus, that may be present in a biological
XX sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX birds and culex mosquitoes, with humans and horses serving as incidental
XX hosts. Infection of humans can lead to meningitis or encephalitis. The
XX invention may allow for accurate and efficient high throughput screening.
XX The present sequence is that of an oligonucleotide probe which is related
XX to the invention.
XX
XX Sequence 51 BP; 18 A; 12 C; 8 G; 13 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.9;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAAATCAAG 22
Db 30 AGCCCTCTTCAGTCCAAATCAAG 51

RESULT 6
ADN36714
ID ADN36714 standard; DNA; 51 BP.
XX
XX ADN36714;
XX
XX 15-JUL-2004 (first entry)
XX
XX West Nile virus detection-related oligonucleotide probe SeqID36.
XX
XX hybridisation assay probe; nucleic acid detection;
XX target-complementary sequence; flavivirus; West Nile virus; WNV;
XX RNA virus; infection; meningitis; encephalitis;
XX high throughput screening; probe; ss.
XX
XX West Nile virus.
XX Enterobacteria phage T7.
XX
XX Key Location/Qualifiers
XX misc_feature 1..27
XX /tag= a
XX /note= "T7 promoter sequence"
XX misc_feature 28..51
XX /tag= b
XX /note= "WNV-complimentary sequence"
XX
XX WO2004036190-A2.
XX
XX 29-APR-2004.
XX

PD 29-APR-2004.
XX
XX 10-OCT-2003; 2003WO-US033639.
XX
XX 16-OCT-2002; 2002US-0418891P.
XX
XX 25-NOV-2002; 2002US-0429006P.
XX
XX 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
XX
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
XX bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Disclosure; SEQ ID NO 38; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
XX detecting a nucleic acid, which is a probe sequence that comprises a
XX target-complementary sequence of bases, and optionally one or more base
XX sequences that are not complementary to the nucleic acid that is to be
XX detected. The hybridisation assay probes and the kits are useful in
XX detecting and amplifying a target nucleic acid sequence, for example
XX flavivirus like West Nile virus, that may be present in a biological
XX sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX birds and culex mosquitoes, with humans and horses serving as incidental
XX hosts. Infection of humans can lead to meningitis or encephalitis. The
XX invention may allow for accurate and efficient high throughput screening.
XX The present sequence is that of an oligonucleotide probe which is related
XX to the invention.
XX
XX Sequence 51 BP; 18 A; 12 C; 8 G; 13 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.9;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAAATCAAG 22
Db 30 AGCCCTCTTCAGTCCAAATCAAG 51

RESULT 7
ADN36715
ID ADN36715 standard; DNA; 51 BP.
XX
XX ADN36715;
XX
XX 15-JUL-2004 (first entry)
XX
XX West Nile virus detection-related oligonucleotide probe SeqID37.
XX
XX hybridisation assay probe; nucleic acid detection;
XX target-complementary sequence; flavivirus; West Nile virus; WNV;
XX RNA virus; infection; meningitis; encephalitis;
XX high throughput screening; probe; ss.
XX
XX West Nile virus.
XX Enterobacteria phage T7.
XX
XX Key Location/Qualifiers
XX misc_feature 1..27
XX /tag= a
XX /note= "T7 promoter sequence"
XX misc_feature 28..51
XX /tag= b
XX /note= "WNV-complimentary sequence"
XX
XX WO2004036190-A2.
XX
XX 29-APR-2004.
XX
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PF 10-OCT-2003; 2003WO-US033639.
XX
XX 16-OCT-2002; 2002US-0418891P.
PR 25-NOV-2002; 2002US-0429006P.
PR 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
XX
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
XX bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Disclosure; SEQ ID NO 37; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
XX detecting a nucleic acid, which is a probe sequence that comprises a
XX target-complementary sequence of bases, and optionally one or more base
XX sequences that are not complementary to the nucleic acid that is to be
XX detected. The hybridisation assay probes and the kits are useful in
XX detecting and amplifying a target nucleic acid sequence, for example
XX flavivirus like West Nile virus, that may be present in a biological
XX sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX birds and culex mosquitoes, with humans and horses serving as incidental
XX hosts. Infection of humans can lead to meningitis or encephalitis. The
XX invention may allow for accurate and efficient high throughput screening.
XX The present sequence is that of an oligonucleotide probe which is related
XX to the invention.
XX
XX Sequence 51 BP; 17 A; 12 C; 9 G; 13 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.9; Indels 0; Gaps 0;
Matches 22; Conservative 0; Mismatches 0;
Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 29 AGCCCTCTTCAGTCCAATCAAG 50
RESULT 8
ADN36694
ID ADN36694 standard; DNA; 69 BP.
AC ADN36694;
XX
XX 15-JUL-2004 (first entry)
XX
XX West Nile virus detection-related oligonucleotide probe SeqID16.
XX
XX hybridisation assay probe; nucleic acid detection;
XX target-complementary sequence; flavivirus; West Nile virus; WNV;
XX RNA virus; infection; meningitis; encephalitis;
XX high throughput screening; probe; ss.
XX
XX West Nile virus.
XX
XX WO2004036190-A2.
XX
XX 29-APR-2004.
XX
XX 10-OCT-2003; 2003WO-US033639.
XX
XX 16-OCT-2002; 2002US-0418891P.
PR 25-NOV-2002; 2002US-0429006P.
PR 24-FEB-2003; 2003US-0449810P.
XX
XX (GENP-) GEN-PROBE INC.
XX
XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX
XX WPI; 2004-389590/36.
XX
XX New hybridization assay probe comprising target-complementary sequence of
XX bases, useful in detecting flavivirus, e.g. West Nile virus.
XX
XX Disclosure; SEQ ID NO 37; 135pp; English.
XX
XX This invention relates to a novel hybridisation assay probe, for
XX detecting a nucleic acid, which is a probe sequence that comprises a
XX target-complementary sequence of bases, and optionally one or more base
XX sequences that are not complementary to the nucleic acid that is to be
XX detected. The hybridisation assay probes and the kits are useful in
XX detecting and amplifying a target nucleic acid sequence, for example
XX flavivirus like West Nile virus, that may be present in a biological
XX sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX birds and culex mosquitoes, with humans and horses serving as incidental
XX hosts. Infection of humans can lead to meningitis or encephalitis. The
XX invention may allow for accurate and efficient high throughput screening.
XX The present sequence is that of an oligonucleotide probe which is related
XX to the invention.
XX
XX Sequence 51 BP; 17 A; 12 C; 9 G; 13 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 22; DB 12; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.9; Indels 0; Gaps 0;
Matches 22; Conservative 0; Mismatches 0;
Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 29 AGCCCTCTTCAGTCCAATCAAG 50
RESULT 9
ABK51710/c
ID ABK51710 standard; cDNA; 365 BP.
XX
XX ABK51710;
XX
XX 27-AUG-2002 (first entry)
XX
XX Partial cDNA for west nile virus capsid protein.
XX
XX Human; ss; IgG leader sequence; west nile virus capsid protein;
XX RNA secondary structure; free energy; gene therapy; cancer;
XX hyperproliferative disease; autoimmune disease; rheumatoid arthritis;
XX multiple sclerosis; Sjogren's syndrome; sarcoidosis; scleroderma;
XX insulin-dependent diabetes mellitus; autoimmune thyroiditis; psoriasis;
XX reactive arthritis; ankylosing spondylitis; polymyositis; vasculitis;
XX dermatomyositis; Crohn's disease; ulcerative colitis.
XX
XX West Nile virus.
XX
XX OS
XX
XX WO200229088-A2.
XX
XX 11-APR-2002.
XX
XX 04-OCT-2001; 2001WO-US031451.
XX
XX 04-OCT-2000; 2000US-0237885P.
XX
XX (UYPE-) UNIV PENNSYLVANIA.
XX
XX Weiner DB, Yang J;
XX
XX WPI; 2002-416682/44.
XX
XX Producing recombinant protein for preparing pharmaceutical compounds to
XX treat, e.g., cancers or autoimmune disorders, comprises predicting
XX secondary structure (SS) of mRNA and modifying DNA to give mRNA with SS
XX having increased free energy.
XX
XX Example 2; Fig 1; 48pp; English.
XX
XX

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CC The invention relates to producing (M1) a protein (I) in a recombinant
 CC expression system (II) comprising: (a) predicting the secondary structure
 CC of mRNA; (b) modifying the native heterologous DNA sequence where the
 CC mRNA transcribed from the modified DNA has a secondary structure with
 CC increased free energy; and (c) using the modified DNA in (II) for
 CC production of (I). Also included are (1) an injectable pharmaceutical
 CC composition comprising a nucleic acid molecule that includes a modified
 CC coding sequence (IV) encoding a protein operably linked to regulatory
 CC elements, where (IV) comprises a higher A/T or A/U content relative to the
 CC A/T or A/U content of the native coding sequence and further comprising a
 CC pharmaceutical carrier and (2) a recombinant viral vector comprising a
 CC nucleic acid molecule that includes (IV). The method is used for
 CC producing a protein in a recombinant expression system. Use of a nucleic
 CC acid or recombinant viral vector that have modified DNA sequences to
 CC improve protein production can be used in gene therapy and for the
 CC treatment of cancers, hyperproliferative diseases, and autoimmune
 CC diseases such as rheumatoid arthritis, multiple sclerosis, Sjogren's
 CC syndrome, sarcoidosis, insulin-dependent diabetes mellitus, autoimmune
 CC thyroiditis, reactive arthritis, ankylosing spondylitis, scleroderma,
 CC polymyositis, dermatomyositis, psoriasis, vasculitis, Crohn's disease and
 CC ulcerative colitis. The present sequence is a cDNA for West Nile virus
 CC capsid protein. Fusion constructs of modified mRNA for the capsid protein
 CC and human IgE leader sequence are used in an experiment to minimise the
 CC free energy of the capsid protein mRNA. Note: The present sequence is not
 CC shown in the specification but was created using the information in
 CC figure 1 and the sequence appearing as ABK51708

XX Sequence 365 BP; 103 A; 80 C; 109 G; 73 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 6; Length 365;

Best Local Similarity 100.0%; Pred. No. 1.2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAAATCAAG 22

Db 95 AGCCCTCTTCAGTCCAAATCAAG 74

RESULT 10

ABQ76684/c

ID ABQ76684 standard; DNA; 366 BP.

XX AC ABQ76684;

DT 13-MAY-2003 (first entry)

DE WNVcwt DNA fragment.

XX Capsid protein; WNVcwt; mRNA secondary structure; cancer;
 KW immunosuppressive; antirheumatic; cytostatic; antiulcer; neuroprotective;
 KW antiarthritic; antidiabetic; antichyroid; antipsoriatic; virucide; gene;
 KW antiparasitic; antiallergic; gene therapy; allergen; multiple sclerosis;
 KW protective immune response; hyperproliferative cell; ulcerative colitis;
 KW hyperproliferative disease; psoriasis; autoimmune disease; psoriasis;
 KW rheumatoid arthritis; Sjogren's syndrome; autoimmune thyroiditis;
 KW insulin dependent diabetes mellitus; Crohn's disease; ds.

OS West Nile virus.

XX Key Location/Qualifiers

FT CDS 1..366

FT /tag= a

FT /product= "WNVcwt"

FT /note= "no start or stop codon given"

XX US2002123099-A1.

XX 05-SEP-2002.

XX 04-OCT-2001; 2001US-00971806.

XX 04-OCT-2000; 2000US-0237885P.

XX

PA (WEIN/) WEINER D B.

PA (YANG/) YANG J.

PI Weiner DB, Yang J;

XX WPI; 2003-066795/06.

XX P-PSDB; ABG73556.

XX Producing protein in recombinant expression system involves predicting
 PT secondary structure of RNA encoding a protein and increasing free energy
 PT for the secondary structure by modifying sequence of DNA encoding the
 PT RNA.

XX Example 2; Fig 1; 25pp; English.

XX This invention describes a novel method for producing a protein by
 CC translation of mRNA from heterologous DNA sequences. The method involves
 CC predicting the secondary structure of mRNA transcribed from a native
 CC heterologous DNA sequence, modifying the sequence where mRNA transcribed
 CC from the modified DNA sequence has a secondary structure with increased
 CC free energy compared to mRNA transcribed from native DNA and using
 CC modified heterologous DNA for protein production. The products of the
 CC invention have immunosuppressive, antineumatic, cytostatic, antiulcer,
 CC neuroprotective, antidiabetic, antichyroid, antipsoriatic, antiparasitic,
 CC virucide, antiparasitic and antiallergic activity and can be used for
 CC gene therapy. The method described is useful for producing a protein in a
 CC recombinant expression system, preferably a cell free in vitro
 CC transcription and translation system, an in vitro cell expression system,
 CC a DNA construct used in direct DNA injection, or a recombinant vector for
 CC delivery of DNA to an individual. The products of the invention are
 CC useful for eliciting broad immune responses against a target protein,
 CC i.e. proteins specifically associated with pathogens such as viruses,
 CC parasites, allergens, or the individual's own abnormal cells.
 CC Compositions containing the products of the invention confer a broad
 CC based protective immune response against hyperproliferative cells that
 CC are characteristic in hyperproliferative diseases including all forms of
 CC cancer and psoriasis. Such compositions are also useful for treating
 CC individuals suffering from autoimmune diseases including rheumatoid
 CC arthritis, multiple sclerosis, Sjogren's syndrome, insulin dependent
 CC diabetes mellitus, autoimmune thyroiditis, Crohn's disease, ulcerative
 CC colitis and psoriasis. This sequence encodes the West Nile virus wild-
 CC type capsid protein described as WNVcwt in the disclosure of the
 CC invention

XX Sequence 366 BP; 103 A; 81 C; 108 G; 74 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 8; Length 366;

Best Local Similarity 100.0%; Pred. No. 1.2;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAAATCAAG 22

Db 96 AGCCCTCTTCAGTCCAAATCAAG 75

RESULT 11

ADQ30647/c

ID ADQ30647 standard; DNA; 967 BP.

XX AC ADQ30647;

XX 23-SEP-2004 (first entry)

XX West Nile virus internal diagnosis control sequence.

XX ss; internal control; West Nile Virus; diagnosis.

XX West Nile virus.

XX WO2004055159-A2.

XX 01-JUL-2004.

XX

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PF 05-DEC-2003; 2003WO-US038750.
XX
PR 12-DEC-2002; 2002US-0432850P.
PR 20-JUN-2003; 2003US-0480431P.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Shyamala V;
XX
XX WPI; 2004-488058/46.
XX
XX New isolated oligonucleotides for accurately diagnosing West Nile virus
PT infection or for capturing, detecting and quantitating West Nile virus in
PT blood samples.
XX
XX Claim 27; SEQ ID NO 17; 56pp; English.
XX
XX The invention relates to an isolated oligonucleotide not more than 60
CC nucleotides in length comprising a nucleotide sequence (S1) of at least
CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
CC 20, 21 or 23 bp) given in the specification derived from the West Nile
CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
CC identity to the nucleotide sequence of (S1), or complements of (S1) and
CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
CC end and/or the 3'-end. The detectable label is a fluorescent label
CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
CC composition and methods are useful for accurately diagnosing West Nile
CC virus infection or for capturing, detecting and quantitating West Nile
CC virus in biological samples, particularly blood samples. This sequence
CC corresponds to an internal control sequence for the detection of WNV
CC sequences using the oligonucleotides of the invention.
XX
XX Sequence 967 BP; 273 A; 206 C; 272 G; 216 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 967;
Best Local Similarity 100.0%; Pred. No. 1.4;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 197 AGCCCTCTTCAGTCCAATCAAG 176

RESULT 12
ADR32078/c
ID ADR32078 standard; DNA; 10945 BP.
XX
XX ADR32078;
XX
XX 18-NOV-2004 (first entry)
XX
XX Genomic DNA of a West Nile virus.
XX
XX analysis; target; real time PCR; ds; genomic.
XX
XX West Nile virus.
XX
XX WO2004072230-A2.
XX
XX 26-AUG-2004.
XX
XX 10-FEB-2004; 2004WO-US002012.
XX
XX 10-FEB-2003; 2003US-00361004.
XX
XX (CLEA-) CLEARANT INC.
XX
XX Mckenney K, Gillmeister L, Marlowe K, Armistead D;
XX
XX WPI; 2004-625843/60.
XX
XX Analyzing a target nucleic acid sequence in a biological material by real

PT time PCR using nucleic acid primers that are separated by at least 750
XX nucleic acid residues in the target sequence.
XX
XX Disclosure; SEQ ID NO 5; 96pp; English.
XX
XX The invention relates to a novel method for analysing a target nucleic
CC acid sequence in a biological material. The method comprises adding at
CC least two nucleic acid primers that hybridise under stringent conditions
CC to predetermined nucleic acid sequences of the target nucleic acid
CC sequence that are separated by at least 750 nucleic acid residues,
CC amplifying the target nucleic acid sequence by PCR, and detecting and
CC quantifying the target nucleic acid sequence. The methods and
CC compositions of the present invention are useful for analysing a target
CC nucleic acid sequence in a biological material by real time PCR using
CC nucleic acid primers that are separated by at least 750 nucleic acid
CC residues in the target sequence. This polynucleotide sequence represents
CC the genomic DNA of a West Nile virus used in the target analysis method
CC of the invention.
XX
XX Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 13; Length 10945;
Best Local Similarity 100.0%; Pred. No. 1.9;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 153 AGCCCTCTTCAGTCCAATCAAG 132

RESULT 13
ADR67768/c
ID ADR67768 standard; DNA; 10945 BP.
XX
XX ADR67768;
XX
XX 18-NOV-2004 (first entry)
XX
XX West Nile virus DNA detected by novel detection method.
XX
XX ds; detection; pathogen.
XX
XX West Nile virus.
XX
XX WO2004072231-A2.
XX
XX 26-AUG-2004.
XX
XX 10-FEB-2004; 2004WO-US002013.
XX
XX 10-FEB-2003; 2003US-00361002.
XX
XX (CLEA-) CLEARANT INC.
XX
XX Mckenney K, Gillmeister L, Marlowe K, Armistead D;
XX
XX WPI; 2004-625844/60.
XX
XX Determining level of potentially active biological pathogens in
PT biological material, by adding nucleic acid primer pairs to biological
PT material, amplifying target nucleic acid by PCR, detecting and
PT quantifying target nucleic acid.
XX
XX Disclosure; SEQ ID NO 5; 111pp; English.
XX
XX The invention relates to a method of determining (MI) level of
CC potentially active biological pathogens in biological material, involves
CC adding at least two nucleic acid primer pairs to biological material,
CC amplifying target nucleic acid sequences by PCR, and detecting and
CC quantifying target nucleic acid sequences, where quantity of the nucleic
CC acid sequences is proportional to number of biological pathogens in
CC biological material. (MI) is useful for determining level of potentially
CC active biological pathogens in a biological material such as cells,

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CC tissues, blood or blood components, proteins, enzymes, immunoglobulins,
 CC botanicals, food, ligaments, tendons, nerves, bone, teeth, skin grafts,
 CC bone marrow, heart valves, cartilage, corneas, arteries, veins, organs,
 CC lipids, carbohydrates, collagen, chitin and its derivatives, forensic
 CC samples, mummified material, human or animal remains, stem cells, islet
 CC of langerhans cells, cells for transplantation, red blood cells, white
 CC blood cells or platelets. The biological pathogen is chosen from
 CC bacteria, viruses, fungi and single cell parasites. The biological
 CC pathogen is chosen from Aspergillus, Candida, Histoplasma, Bacillus,
 CC Saccharomyces, Clostridioides, Cryptococcus, Escherichia, Bacillus,
 CC Campylobacter, Helicobacter, Listeria, Clostridium, Streptococcus,
 CC Enterococcus, Staphylococcus, Brucella, Haemophilus, Salmonella,
 CC Yersinia, Pseudomonas, Serratia, Enterobacter, Klebsiella, Proteus,
 CC Citrobacter, Corynebacterium, Propionibacterium and Coxiella. The
 CC biological pathogen is chosen from Adeno-associated virus (AAV),
 CC California encephalitis virus, Coronavirus, Coxsackievirus-A,
 CC Coxsackievirus-B, Eastern equine encephalitis virus (EEEV), Echovirus,
 CC Hantavirus, Hepatitis A virus (HAV), Hepatitis C virus (HCV), Hepatitis
 CC delta virus (HDV), Hepatitis E virus (HEV), Hepatitis G virus (HGV), HIV,
 CC Human T-lymphotrophic virus (HTLV), Influenza virus (Flu virus), Measles
 CC virus (Rubella), Mumps virus, Norwalk virus, Parainfluenza virus, Polio
 CC virus, Rabies virus, Respiratory Syncytial virus, Rhinovirus, Rubella
 CC virus, Saint Louis encephalitis virus, Western equine encephalitis virus
 CC (WEEV), Yellow fever virus, Adenovirus, Cytomegalovirus (CMV), Epstein-
 CC Barr virus (EBV), Hepatitis B virus (HBV), Herpes simplex virus 1, Herpes
 CC simplex virus 2, Molluscum contagiosum, Papilloma virus (HPV), Smallpox
 CC virus (Variola), Vaccinia virus, Venezuelan equine encephalitis virus
 CC (VEEV), Ebola virus, West Nile virus, Human Parvovirus B19 and Rotavirus.
 CC (M1) is useful for determining the effectiveness of a sterilization
 CC process applied to a biological material. (M1) is useful in determining
 CC whether the biological pathogen is inactive or active. (M1) enables
 CC determination of whether the particular biological pathogen is present in
 CC a biological material as shown by amplification of first target sequence
 CC and whether the biological pathogen is inactive or active. (M1) enables
 CC evaluation of the effectiveness of sterilization processes, and
 CC determination of both the original level and the residual level of
 CC potentially active biological pathogens. This sequence corresponds to a
 CC West Nile virus DNA detected by the method of the invention.

XX Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 13; Length 10945;
 Best Local Similarity 100.0%; Pred. No. 1.9;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
 Db 153 AGCCCTCTTCAGTCCAATCAAG 132

RESULT 14
 ADN98022/c
 ID ADN98022 standard; DNA; 10975 BP.

XX AC ADN98022;

XX DT 29-JUL-2004 (first entry)

XX DE West Nile Virus isolate 2741 complete genome sequence.

XX ds; West Nile Virus; envelope protein; glycoprotein E; flavivirus;
 XX Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.

OS West Nile virus.

XX WO2004040263-A2.

XX PD 13-MAY-2004.

XX PF 31-OCT-2003; 2003WO-US034823.

XX PR 31-OCT-2002; 2002US-0422755P.

XX PR 06-JUN-2003; 2003US-0476513P.

XX (HEAL-) HEALTH RES INC.
 XX PA Wong SJ, Pei-Yong S;
 XX PI WPI; 2004-400223/37.
 XX DR GENBANK; AF206518.

XX New diagnostic kit comprising West Nile Virus (WNV) envelope protein
 PT reactive with antibody against WNV and cross-reactive with antibody
 PT against a flavivirus, useful in diagnosing flavivirus infection caused by
 PT DENV, WNV, JEV or SLEV.

PS Disclosure; Fig 37; 212pp; English.

XX The invention relates to a diagnostic kit comprising at least one
 CC isolated and purified polypeptide comprising a West Nile Virus (WNV)
 CC envelope (E) protein or its immunogenic fragment having a native
 CC conformation or non-denatured structure and that is reactive with
 CC antibodies against WNV and cross-reactive with antibodies against a
 CC flavivirus. The diagnostic kit is useful in diagnosing flavivirus
 CC infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to
 CC the complete nucleotide sequence of the WNV isolate 2741.

XX Sequence 10975 BP; 3007 A; 2460 C; 3149 G; 2359 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 10975;
 Best Local Similarity 100.0%; Pred. No. 2;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
 Db 177 AGCCCTCTTCAGTCCAATCAAG 156

RESULT 15
 ABZ68481/c
 ID ABZ68481 standard; DNA; 11029 BP.

XX AC ABZ68481;

XX DT 22-APR-2003 (first entry)

XX DE Nucleotide sequence of the genome of West Nile virus IS-98-ST1.

XX WNV; IS-98-ST1; Flavivirus; infection; encephalitis; gene; ss.

XX OS West nile virus.

XX Key Location/Qualifiers
 FT 97-10397
 FT CDS /*tag= a
 FT /product= "polypotein"

XX WO200281511-A1.

XX PD 17-OCT-2002.

XX PF 04-APR-2002; 2002WO-FR001168.

XX PR 04-APR-2001; 2001FR-00004599.

XX PR 06-SEP-2001; 2001FR-00011525.

XX (INSP) INST PASTEUR.

XX (KIMR-) KIMRON VETERINARY INST.

XX Despres P, Deubel V, Guenet J, Drouet M, Malkinson M, Banet C;
 XX Frenkiel M, Courageot M, Coulibaly F, Cateau A, Flamand M, Weber P;
 XX Ceccaldi P;

XX WPI; 2003-058498/05.

XX P-PSDB; ABP70647.

PT New neurovirulent strain of West Nile virus, useful in diagnosis and
 PT screening for antiviral agents, also related nucleic acids, proteins and
 PT antibodies.

XX Claim 1; Page 34-49; 68pp; French.

XX The present sequence represents the genome of a strain of West Nile virus
 CC (WNV), designated IS-98-ST1. This strain is a neuroinvasive and
 CC neurovirulent strain of WNV. Polynucleotides and polypeptides derived
 CC from the IS-98-ST1 genome are useful for diagnosis and prognosis of
 CC Flavivirus infection, specifically WNV-mediated encephalitis. They are
 CC also useful to raise specific antibodies, for recombinant expression of
 CC WNV proteins or peptides (for diagnosis, production of antibodies and
 CC identification of specific binding partners in cells), for identifying
 CC cellular genes implicated in resistance to viral infection, and for
 CC screening for anti-Flavivirus agents

XX Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 8; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 2;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22

Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 16

ABV74821/C

ID ABV74821 standard; DNA; 11029 BP.

XX AC ABV74821;

XX 28-MAR-2003 (first entry)

XX West Nile virus strain NY99-flamingo 382-99 complete genome.

XX Virucide; hepatotropic; antiinflammatory; antiviral; OAS;

XX 2'-5'-oligoadenylate synthase; Flavivirus infection; gene; ss.

XX West Nile Virus.

XX Key Location/Qualifiers

XX 97. .10398

XX /*tag= a

XX /product= "West Nile Virus protein"

XX WO2002081741-A2.

XX 17-OCT-2002.

XX 04-APR-2002; 2002WO-FR001169.

XX 04-APR-2001; 2001FR-00004598.

XX (INSP) INST PASTEUR.

XX (CNRS) CNRS CENT NAT RECH SCI.

XX Guenet J, Mashimo T, Simon-Chazottes D, Montagutelli X;

XX Frankiel M, Despres P, Deubel V, Bonhomme F, Lucas M;

XX WPI; 2003-058566/05.

XX P-PSDB; ABB98821.

XX Identifying stimulators of oligoadenylate synthase family genes, useful
 PT as antiviral agents against Flavivirus, also mutated genes responsible
 PT for sensitivity to virus.

XX Example 1; Page 52-67; 93pp; French.

XX The present invention relates to a method for identifying compounds (I)
 CC that can stimulate a gene of the OAS (2'-5'-oligoadenylate synthase)

CC family. The method comprises: (a) inducing expression of the OAS gene in
 CC a culture of cells from a non-human mammal (Flvr/Flvr or Flvr/Flvs;
 CC indicating resistance or sensitivity to Flavivirus infection); (b)
 CC treating cells with test compound; and (c) measuring activity of OAS gene
 CC relative to a control. (I) are potentially useful as antiviral agents for
 CC treating infections by Flaviviruses (e.g. hepatitis C; dengue; yellow
 CC fever and various forms of encephalitis). Genomic OAS DNA and derived
 CC cDNA, also the encoded proteins, are useful: (a) for treating Flavivirus
 CC infection; (b) in screening for anti-flavivirus agents, and (c) for
 CC evaluating sensitivity of subjects to Flavivirus infection and their
 CC likely response to interferon treatment, e.g. to identify patients at
 CC risk of developing severe forms of such infections. The present sequence
 CC is West Nile Virus strain NY99-flamingo 382-99 (IS-98-ST1) complete
 CC genome, which was used in an example from the invention. West Nile Virus
 CC is one such Flavivirus

SQ Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 10; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 2;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22

Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 17

ADN98023/C

ID ADN98023 standard; DNA; 11029 BP.

XX AC ADN98023;

XX 29-JUL-2004 (first entry)

XX West Nile Virus isolate 3356 complete genome sequence.

XX ds; West Nile Virus; envelope protein; glycoprotein E; flavivirus;

XX Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.

XX West Nile virus.

XX WO2004040263-A2.

XX 13-MAY-2004.

XX 31-OCT-2003; 2003WO-US034823.

XX 31-OCT-2002; 2002US-0422755P.

XX 06-JUN-2003; 2003US-0476513P.

XX (HEAL-) HEALTH RES INC.

XX Wong SJ, Pei-Yong S;

XX WPI; 2004-400223/37.

XX GENBANK; AF404756.

XX New diagnostic kit comprising West Nile Virus (WNV) envelope protein
 PT reactive with antibody against WNV and cross-reactive with antibody
 PT against a flavivirus, useful in diagnosing flavivirus infection caused by
 PT DENV, WNV, JEV or SLEV.

XX Disclosure; Fig 38; 212pp; English.

XX The invention relates to a diagnostic kit comprising at least one
 CC isolated and purified polypeptide comprising a West Nile Virus (WNV)
 CC envelope (E) protein or its immunogenic fragment having a native
 CC conformation or non-denatured structure and that is reactive with
 CC antibodies against WNV and cross-reactive with antibodies against a
 CC flavivirus. The diagnostic kit is useful in diagnosing flavivirus
 CC infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to
 CC the complete nucleotide sequence of the WNV isolate 3356.

```
XX SQ Sequence 11029 BP; 3017 A; 2466 C; 3172 G; 2374 T; 0 U; 0 Other;
Query Match 100.0%; Score 22; DB 12; Length 11029;
Best Local Similarity 100.0%; Pred. No. 2;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 18
ADN36705
ID ADN36705 standard; DNA; 22 BP.
XX AC ADN36705;
XX DT 15-JUL-2004 (first entry)
XX DE West Nile virus detection-related PCR primer SeqID27.
XX KW hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; PCR; primer; ss.
XX OS West Nile virus.
XX PN WO2004036190-A2.
XX PD 29-APR-2004.
XX PF 10-OCT-2003; 2003WO-US033639.
XX PR 16-OCT-2002; 2002US-0418891P.
XX PR 25-NOV-2002; 2002US-0429006P.
XX PR 24-FEB-2003; 2003US-0449810P.
XX PA (GENP-) GEN-PROBE INC.
XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX DR
XX PT New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX PS Example 2; SEQ ID NO 27; 135pp; English.
XX CC This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of a PCR primer which is related to the
CC invention.
XX SQ Sequence 22 BP; 6 A; 8 C; 2 G; 6 T; 0 U; 0 Other;
Query Match 95.5%; Score 21; DB 12; Length 22;
Best Local Similarity 100.0%; Pred. No. 2.4;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAA 21
Db 2 AGCCCTCTTCAGTCCAATCAA 22
```

```
RESULT 19
ADN36717
ID ADN36717 standard; DNA; 49 BP.
XX AC ADN36717;
XX DT 15-JUL-2004 (first entry)
XX DE West Nile virus detection-related oligonucleotide probe SeqID39.
XX KW hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.
XX OS West Nile virus.
XX OS Enterobacteria phage T7.
XX FH Key Location/Qualifiers
FH misc_feature 1..27
FT /tag= a
FT /note= "T7 promoter sequence"
FT misc_feature 28..49
FT /tag= b
FT /note= "WNV-complimentary sequence"
XX PN WO2004036190-A2.
XX PD 29-APR-2004.
XX PF 10-OCT-2003; 2003WO-US033639.
XX PR 16-OCT-2002; 2002US-0418891P.
XX PR 25-NOV-2002; 2002US-0429006P.
XX PR 24-FEB-2003; 2003US-0449810P.
XX PA (GENP-) GEN-PROBE INC.
XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX DR
XX PT New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.
XX PS Disclosure; SEQ ID NO 39; 135pp; English.
XX CC This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.
XX SQ Sequence 49 BP; 17 A; 12 C; 7 G; 13 T; 0 U; 0 Other;
Query Match 95.5%; Score 21; DB 12; Length 49;
Best Local Similarity 100.0%; Pred. No. 2.7;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAA 21
Db 29 AGCCCTCTTCAGTCCAATCAA 49
```


[illegible]

KW hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; PCR; primer; ss.

XX
 OS West Nile virus.
 OS Enterobacteria phage T7.
 XX
 FH Key Location/Qualifiers
 FT misc_feature 1..27
 FT /*tag= a
 FT /note= "T7 promoter sequence"
 FT 28..51
 FT /*tag= b
 FT /note= "WNV-complimentary sequence"
 XX

XX WO2004036190-A2.

XX 29-APR-2004.

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

XX 25-NOV-2002; 2002US-0429006P.

XX 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 XX Example 6; SEQ ID NO 40; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of a PCR primer which is related to the
 CC invention.

XX Sequence 51 BP; 18 A; 13 C; 7 G; 13 T; 0 U; 0 Other;

Query Match 90.9%; Score 20; DB 12; Length 51;
 Best Local Similarity 100.0%; Pred. No. 8.3;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCAATCA 20

DB 32 AGCCCTCTTCAGTCCAATCA 51

RESULT 23

ADN36701

ID ADN36701 standard; DNA; 24 BP.

XX AC ADN36701;

XX 15-JUL-2004 (first entry)

DE West Nile virus detection-related PCR primer SeqID23.

XX hybridisation assay probe; nucleic acid detection;

KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 XX high throughput screening; PCR; primer; ss.

OS West Nile virus.

XX WO2004036190-A2.

XX 29-APR-2004.

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

XX 25-NOV-2002; 2002US-0429006P.

XX 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
 XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.
 XX Example 2; SEQ ID NO 23; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of a PCR primer which is related to the
 CC invention.

XX Sequence 24 BP; 8 A; 8 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 86.4%; Score 19; DB 12; Length 24;
 Best Local Similarity 100.0%; Pred. No. 23;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 CCTCTTCAGTCCAATCAAG 22

DB 1 CCTCTTCAGTCCAATCAAG 19

RESULT 24

ADK13681/c

ID ADK13681 standard; DNA; 10962 BP.

XX AC ADK13681;

XX 20-MAY-2004 (first entry)

XX West Nile Virus DNA sequence, SEQ ID 1.

XX Virucide; Immunostimulant; flavivirus;
 KW envelope protein domain III polypeptide; envelope protein; gene; ss.

XX West Nile virus.

XX Key Location/Qualifiers

XX 97..10389

XX /*tag= a

XX /product= "West Nile Virus protein"

XX WO2004016586-A2.

XX PD 26-FEB-2004.
XX PF 18-AUG-2003; 2003WO-US025681.
XX PR 16-AUG-2002; 2002US-0403893P.
XX PR 08-FEB-2003; 2003US-0445581P.
XX PA (TEXA) UNIV TEXAS SYSTEM.
XX PI Barrett A, Beasley D, Holbrook M;
XX WPI; 2004-203756/19.
DR P-PSDB; ADK13682.
XX Diagnosing flavivirus infection by contacting a sample from a human or
PT animal with a flavivirus envelope protein domain III polypeptide, and
PT detecting formation of an immunocomplex between the envelope protein and
PT antibodies in the sample.
XX Disclosure; SEQ ID NO 1; 110pp; English.
XX
XX The present invention relates to a method for screening for a flavivirus
CC in a subject or animal host. The method comprises: contacting a sample
CC from the subject with a composition comprising a flavivirus envelope
CC protein domain III polypeptide (ADK13683-ADK13701) under conditions that
CC permit formation of specific immunocomplex between an antibody in the
CC sample and the envelope protein domain III polypeptide; and detecting
CC whether a specific immunocomplex is formed. The present sequence is the
CC coding sequence for West Nile Virus protein, from which E protein
CC envelope protein domain III polypeptide (ADK13683) is derived.
XX
XX Sequence 10962 BP; 2997 A; 2497 C; 3100 G; 2368 T; 0 U; 0 Other;
SQ Query Match 85.5%; Score 18.8; DB 12; Length 10962;
Best Local Similarity 90.9%; Pred. No. 68;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
DB 195 AGCCCTCTTAGTCTATCAAG 174

RESULT 25
ADQ97910/c
ID ADQ97910 standard; DNA; 44920 BP.
XX
XX ADQ97910;
XX
XX 07-OCT-2004 (first entry)
XX Human cancer associated sequence HD11-022, SEQ ID 887.
XX
XX Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Human; ds.
XX
XX Homo sapiens.
XX
XX WO2004060304-A2.
XX
XX 22-JUL-2004.
XX
XX 22-DEC-2003; 2003WO-US041389.
XX
XX 27-DEC-2002; 2002US-00330773.
XX
XX (SAGR-) SAGRES DISCOVERY INC.
XX
XX Morris DW, Malandro MS;
XX
XX WPI; 2004-543781/52.
XX
XX New isolated cancer associated nucleic acids comprising at least 10
PT contiguous nucleotides, useful for diagnosing, preventing and/or treating

PT cancers such as leukemia and lymphoma.
XX
XX Claim 1; SEQ ID NO 887; 199pp; English.
XX
XX The present invention relates to cancer associated sequences (ADQ97025-
CC ADQ98004). The sequences are useful for the diagnosis, prevention and/or
CC treatment of cancer, such as leukemia and lymphoma. Note: The sequence
CC data for this patent did not form part of the printed specification, but
CC was obtained in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 44920 BP; 13235 A; 9200 C; 9459 G; 13026 T; 0 U; 0 Other;
SQ Query Match 85.5%; Score 18.8; DB 12; Length 44920;
Best Local Similarity 90.9%; Pred. No. 84;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGCCCTCTTCAGTCCCAATCAAG 22
DB 42318 AGCCCTCTTCAATCTAATCAAG 42297

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Job time : 202.688 secs

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OM nucleic - nucleic search, using sw model

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Scoring table: IDENTITY_NUC
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Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 100 summaries

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EST:*
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3: gb_hc:*
4: gb_est3:*
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6: gb_est5:*
7: gb_est6:*
8: gb_ges1:*
9: gb_ges2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|---------------------|
| 1 | 19 | 86.4 | 1174 | 8 | CC229626 CH261-46H |
| 2 | 18.8 | 85.5 | 767 | 2 | BE541043 601064391 |
| 3 | 18.8 | 85.5 | 872 | 2 | BF687801 602066853 |
| C 4 | 18.4 | 83.6 | 935 | 9 | CL901525 CSRC1000 |
| C 5 | 17.8 | 80.9 | 169 | 1 | AA693379 an21906.s |
| C 6 | 17.8 | 80.9 | 185 | 2 | BF833890 RC1-HT088 |
| 7 | 17.8 | 80.9 | 193 | 2 | AW800621 MR1-UM006 |
| 8 | 17.8 | 80.9 | 205 | 2 | BF111499 7128f04.x |
| C 9 | 17.8 | 80.9 | 270 | 1 | AA884812 am28b05.s |
| C 10 | 17.8 | 80.9 | 270 | 2 | AW819544 RC5-ST029 |
| C 11 | 17.8 | 80.9 | 271 | 1 | AA437133 zv53e10.s |
| 12 | 17.8 | 80.9 | 289 | 2 | BB720494 BB720494 |
| 13 | 17.8 | 80.9 | 295 | 4 | BM151302 TCBAPI061 |
| C 14 | 17.8 | 80.9 | 309 | 1 | AI2033209 qr23h08.x |
| C 15 | 17.8 | 80.9 | 310 | 7 | T46996 yb12b06.r1 |
| 16 | 17.8 | 80.9 | 328 | 1 | AA969466 co81d01.s |
| C 17 | 17.8 | 80.9 | 328 | 2 | AW615159 hg73h02.x |
| C 18 | 17.8 | 80.9 | 339 | 1 | AI307235 tb18c07.x |
| C 19 | 17.8 | 80.9 | 339 | 1 | AI307235 wa89g09.x |
| C 20 | 17.8 | 80.9 | 339 | 1 | AI631503 A1631503 |
| C 21 | 17.8 | 80.9 | 339 | 1 | AI633750 tt28b04.x |
| C 22 | 17.8 | 80.9 | 339 | 1 | AI954798 wq33d07.x |
| C 23 | 17.8 | 80.9 | 339 | 1 | AI954834 wq33h04.x |
| C 24 | 17.8 | 80.9 | 339 | 1 | AI962281 wq46e04.x |

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AA749323 ny12c01.8
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AA768437 ob22f07.8
AA535707 nf88d04.8
AW291203 UI-H-B12-
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A1074659 ox82g07.8
CN545186 EST 17130
A1041258 ov66a02.x
BM998692 UI-H-DT1-
AA417921 zv94b03.8
AA834127 of26g07.8
AA073711 xb01h10.x
A1418470 tg48e05.x
AA768438 ob22f08.8
AA418172 zv94f03.x
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BQ012758 UI-1-BC1p
AI088652 qb14a07.x
AA936255 on75b04.8
AA771206 xv13e09.x
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BQ774467 UI-H-E21-
BE440143 HTM1-954R
AI400162 tg67g09.x
AW128873 xe89c02.x
AA864874 oh03d08.8
BF890876 PM2-MT010
B1962326 ie60d12.y
B1966885 ie62f08.x
B1963087 ie62f08.y
AI088606 qb14e01.x
AV735110 AV735110
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CB267636 1008542 H
AV715652 AV715652
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AL701719 DKF2p686H
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B1966756 ie60d12.x
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BG500991 602546583
BE739605 601556586
B1553318 603193449
BG535175 602562794
BG505886 601859895
BM713190 UI-B-EJ0-
BF213738 601847628
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CE639326 AGENCOURT
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B1562028 603355083
BE566608 601340137

7 T46995
2 AW291932
2 AW237469
8 AZ693999
8 AA749323
1 A1150471
1 AA768437
1 AA535707
2 AW291203
2 A1242291
1 A1074659
7 CN545186
1 A1041258
5 BM998692
4 AA417921
4 AA834127
2 AA073711
2 A1418470
1 AA768438
1 AA418172
5 BQ942517
5 BP305540
6 BY560157
6 BE379976
1 AA159601
2 AW275079
5 BQ012758
1 A1088652
1 AA936255
2 AA771206
5 BX279662
5 BQ774467
2 BE440143
1 A1400162
2 AA128873
2 AA864874
2 BF890876
4 B1962326
4 B1966885
4 B1963087
1 A1088606
1 AV735110
2 AW970416
6 CB267636
1 AV715652
4 BF970233
4 BG723880
1 AL701719
5 BX094624
4 BM670715
1 AI309768
5 BP379530
4 B1966756
6 CB160326
2 BE739363
1 AV716017
7 CV027350
6 BG500991
2 BE739605
4 B1553318
4 BG535175
4 BG505886
4 BM713190
8 BE3640
6 BF213738
6 CD639326
6 CE750532
6 CD641555
2 AV762336
2 BB603221
1 AV758775
4 B1562028
2 BE566608

339 80.9
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766 80.9

C 25
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C 90
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C 92
C 93
C 94
C 95
C 96
C 97

98 17.8 80.9 766 9 BX204503 BX204503 Danio rerio
 99 17.8 80.9 775 2 BF529524 BF529524 602043291
 c 100 17.8 80.9 785 1 AV757859 AV757859 AV757859

ALIGNMENTS

RESULT 1
 CC229626 1174 bp DNA linear GSS 12-MAY-2003
 LOCUS CH261-46H2 Sp6.1 CH261 Gallus gallus genomic clone CH261-46H2,
 DEFINITION genomic survey sequence.
 ACCESSION CC229626
 VERSION CC229626
 KEYWORDS CC229626.1 GI:30556289
 SOURCE GSS.
 ORGANISM Gallus gallus (chicken)
 Archosauria; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Phasianidae; Gallus.
 REFERENCE 1 (bases 1 to 1174)
 AUTHORS Krenitzki,C., Higginbotham,J., Wylie,K., Carter,J., McPherson,J.,
 Warren,W., Graves,T., Mardis,E. and Wilson,R.
 TITLE Gallus gallus BAC End Reads
 JOURNAL Unpublished (2003)
 COMMENT Contact: Richard K. Wilson
 Genome Sequencing Center
 Washington University School of Medicine
 Email: submissions@wustl.edu
 Insert Length: 182000 Std Error: 0.00
 Seq primer: Sp6 ATTTAGGTGACACTATAG
 Class: BAC ends
 High quality sequence start: 329
 High quality sequence stop: 482.
 Location/Qualifiers
 1..1174
 /organism="Gallus gallus"
 /mol_type="genomic DNA"
 /strain="Red Jungle Fowl"
 /db_xref="taxon:9031"
 /clone="CH261-46H2"
 /sex="female"
 /cell_line="UCD001, inbred 256"
 /clone_lib="CH261"
 /note="Vector: pTARBAC2.1; Site 1: EcoRI; Site 2: EcoRI;
 CH261 Female Chicken library - for library and clone
 ordering information: http://www.chori.org/bacpac"

Query Match 86.4%; Score 19; DB 8; Length 1174;
 Best Local Similarity 100.0%; Pred. No. 3.8e+02;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 4 CCTCTTCAGTCCCAATCAAG 22
 |||||
 Db 399 CCTCTTCAGTCCCAATCAAG 417

ORIGIN

Query Match 86.4%; Score 19; DB 8; Length 1174;
 Best Local Similarity 100.0%; Pred. No. 3.8e+02;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 4 CCTCTTCAGTCCCAATCAAG 22
 |||||
 Db 399 CCTCTTCAGTCCCAATCAAG 417

RESULT 2
 BE541043 767 bp mRNA linear EST 09-AUG-2000
 LOCUS 601064391F1 NIH_MGC_10 Homo sapiens cdna clone IMAGE:3450647 5',
 DEFINITION mRNA sequence.
 ACCESSION BE541043
 VERSION BE541043.1 GI:9769787
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 767)
 AUTHORS NIH-MGC http://mgc.nci.nih.gov/.

TITLE
JOURNAL
COMMENT

National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: ATCC
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: Incyte Genomics, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LHAM8429 row: f column: 24
 High quality sequence stop: 523.
 Location/Qualifiers
 1..767
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:3450647"
 /cell_line="MGC36"
 /lab_host="DH10B"
 /clone_lib="NIH_MGC_10"
 /note="Organ: cervix; Vector: pCMV-SPORT6; Site 1: NotI;
 Site 2: SalI; Cloned unidirectionally. Primer: Oligo dt.
 Average insert size 1.5 kb. Library prepared by Life
 Technologies."

FEATURES
source

Query Match 85.5%; Score 18.8; DB 2; Length 767;
 Best Local Similarity 90.9%; Pred. No. 4.5e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 AGCCTCTTCAGTCCCAATCAAG 22
 |||||
 Db 107 AGTCTCTTCAGTCCCAATCAAG 128

ORIGIN

Query Match 85.5%; Score 18.8; DB 2; Length 767;
 Best Local Similarity 90.9%; Pred. No. 4.5e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 AGCCTCTTCAGTCCCAATCAAG 22
 |||||
 Db 107 AGTCTCTTCAGTCCCAATCAAG 128

RESULT 3
 BF687801 872 bp mRNA linear EST 22-DEC-2000
 LOCUS 602068853F1 NIH_MGC_57 Homo sapiens cdna clone IMAGE:4065901 5',
 DEFINITION mRNA sequence.
 ACCESSION BF687801
 VERSION BF687801.1 GI:11973209
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 872)
 AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: ATCC
 cDNA Library Preparation: CLONETECH Laboratories, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LLCM902 row: j column: 14
 High quality sequence stop: 623.
 Location/Qualifiers
 1..872
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4065901"
 /tissue_type="globlastoma"
 /lab_host="DH10B (T1 phage-resistant)"
 /clone_lib="NIH_MGC_57"

FEATURES
source

Query Match 85.5%; Score 18.8; DB 2; Length 767;
 Best Local Similarity 90.9%; Pred. No. 4.5e+02;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 AGCCTCTTCAGTCCCAATCAAG 22
 |||||
 Db 107 AGTCTCTTCAGTCCCAATCAAG 128

/note="Organ: brain; Vector: pDNR-LIB (Clontech); Site_1: SfiI (ggcgcccgcc); Site_2: SfiI (ggccattatggcc); Double-stranded cDNA was prepared from cell line RNA. 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CAGGCCATTATGCC-3' and 3' adaptor sequence: 5'-ATTCTAGAGCCGAGCGCGGACATG-dT(30)BN-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.55 kb (range 0.9-4.0 kb). 12/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA)."

ORIGIN

Query Match 85.5%; Score 18.8; DB 2; Length 872;
Best Local Similarity 90.9%; Pred. No. 4.5e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAAATCAAG 22

Db 319 AGTCTCTTCAGTCCAAATCAAG 340

RESULT 4

CL901625/c
LOCUS
DEFINITION CL901625 935 bp DNA linear GSS 30-AUG-2004
1639HC06E05, genomic survey sequence.

ACCESSION CL901625

VERSION CL901625.1 GI:51663670

KEYWORDS GSS.

SOURCE Triticum aestivum (bread wheat)

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poideae; Triticeae; Triticum.

1 (bases 1 to 935)

Lamoureux,D., Peterson,D.G., Li,W., Fellers,J.P. and Gill,B.S.

Cot-based cloning and sequencing (CBCS) efficiently removes

sequence repeats and increases gene ratio in bread wheat

Unpublished (2004)

Contact: Gill BS

Department of Plant Pathology

Kansas State University

4024 Throckmorton, Manhattan, KS 66506-5502, USA

Tel: 785 532 1391

Fax: 785 532 5692

Email: bgill@ksu.edu

Seq primer: 17

Class: sheared ends.

FEATURES

source

1. 935
/organism="Triticum aestivum"
/mol_type="genomic DNA"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="1639HC06E05"
/tissue_type="whole plant"
/dev_stage="young shoot"
/clone_lib="1639HC library"

ORIGIN

Query Match 83.6%; Score 18.4; DB 9; Length 935;
Best Local Similarity 95.0%; Pred. No. 7.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 CCCTCTTCAGTCCAAATCAAG 22

Db 216 CCCTCTTCAGTCCAAATCAAG 197

RESULT 5

AA693379/c
LOCUS
DEFINITION AA693379 169 bp mRNA linear EST 12-JAN-1999

DEFINITION

ab21906.g1 Soares_parathyroid_tumor_NbHPA Homo sapiens cDNA clone

1239514.3' similar to TR:Q13227 Q13227 GPS2.1, mRNA sequence.

AA693379

VERSION AA693379.1 GI:2694317

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 169)

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: ccapbe-remail.nih.gov

CNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima

Bonaldo, Ph.D.

CNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality

Possible reversed clone: similarity on wrong strand

Insert Length: 912 Std Error: 0.00

Seq primer: -40m13 fwd. ET from Amersham

High quality sequence stop: 1.

FEATURES

source

1. 169
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="1239514"
/tissue_type="parathyroid tumor"
/dev_stage="adult"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares_parathyroid_tumor_NbHPA"
/note="Organ: parathyroid gland; Vector: pT7T3D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer
[5'-TGTTACCAATCTGAGTGGAGCGCGCCACCAATTTTTTTTTTTTTTTT
TTTTT-3'], double-stranded cDNA was size selected, ligated
to Eco RI adaptors (Pharmacia), digested with Not I and
cloned into the Not I and Eco RI sites of a modified pT7T3
vector (Pharmacia). Library went through one round of
normalization to a Cot = 5. Library constructed by Bento
Soares and M.Fatima Ronaldo. RNA from sporadic parathyroid
adenomas was kindly provided by Dr. Stephen Marx, National
Institute of Diabetes and Digestive and Kidney Diseases,
NIH."

ORIGIN

Query Match 80.9%; Score 17.8; DB 1; Length 169;
Best Local Similarity 90.5%; Pred. No. 1.1e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GCCCTCTTCAGTCCAAATCAAG 22

Db 130 GCCATCTTCAGCCCAATCAAG 110

RESULT 6

BF833890/c

LOCUS

DEFINITION RC1-HT0881-041100-019-b04 HT0881 Homo sapiens cDNA, mRNA sequence.

ACCESSION BF833890

VERSION BF833890.1 GI:12183723

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 185)

REFERENCE AUTHORS

Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V.,
O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.J.

Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags

TITLE

JOURNAL
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

MEDLINE PUBMED

20202663

COMMENT

Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?t1=RC1&t2=RC1-HT0881-
041100-019-b04&t3=2000-11-04&t4=1)

Seq primer: puc 18 forward
High quality sequence stop: 185.
Location/Qualifiers
1..185

FEATURES

source

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="HT0881"

/note="Organ: head neck; Vector: puc18; Site 1: SmaI;
Site 2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the puc 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

ORIGIN

Query Match 80.9%; Score 17.8; DB 2; Length 185;
Best Local Similarity 90.5%; Pred. No. 1.1e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GCCCTCTTCAGTCCCAATCAAG 22

|||||

Db 29 GCCATCTTCAGTCCCAATCCAG 9

RESULT 7

AW800621

LOCUS

MR1-UM0063-080300-001-c08 UM0063 Homo sapiens CDNA, mRNA sequence.

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS

Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V.,
O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.J.

Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags

JOURNAL

MEDLINE

20202663

PUBMED

COMMENT

Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?t1=RC1&t2=MR1-UM0063-080
300-001-c08&t3=2000-03-08&t4=1)

Seq primer: puc 18 forward

High quality sequence start: 14

High quality sequence stop: 193.

FEATURES

source

1..193
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="UM0063"

/note="Organ: uterus; Vector: puc18; Site 1: SmaI; Site 2:
SmaI; A mini-library was made by cloning products derived
from ORESTES PCR (U.S. Letters Patent application No.
196,716 - Ludwig Institute for Cancer Research) profiles
into the puc 18 vector. Reverse transcription of tissue
mRNA and cDNA amplification were performed under low
stringency conditions."

ORIGIN

Query Match 80.9%; Score 17.8; DB 2; Length 193;

Best Local Similarity 90.5%; Pred. No. 1.1e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GCCCTCTTCAGTCCCAATCAAG 22

|||||

Db 98 GCCATCTTCAGTCCCAATCCAG 118

RESULT 8

BF111499

LOCUS

7128f04.x1 Soares NSF_P8_9W_OT_PA_P_S1 Homo sapiens CDNA clone

IMAGE:3522942 3' similar to TR:092478 092478 C-TYPE LECTIN. [1] ;

mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Contact: Robert Strausberg, Ph.D.

Email: cgap@remail.nih.gov

This clone is available royalty-free through LNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Trace considered overall poor quality

Seq primer: -40UP from Gibco

High quality sequence stop: 1.

FEATURES

source

1..205
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"


```

RESULT 11
AA437133/c
LOCUS AA437133.1 Soares_testis_NHT Homo sapiens cDNA clone IMAGE:757386
DEFINITION 3', mRNA sequence.
ACCESSION AA437133
VERSION AA437133.1 GI:2142047
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 271)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J.,
Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B.,
White, Y., Wylie, T., Waterston, R. and Wilson, R.
TITLE WashU-Merck EST Project 1997
JOURNAL Unpublished (1997)
COMMENT Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Seq primer: -41m13 fwd. ET from Amersham
High quality sequence stop: 166.
FEATURES
source
1..271
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:5978000"
/db_xref="taxon:9606"
/clone="IMAGE:757386"
/sex="male"
/lab_host="DH10B"
/clone_lib="Soares testis NHT"
/note="Vector: pTT3D-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was prepared from mRNA obtained from Clontech
Laboratories, Inc., and primed with a Not I - oligo(dT)
primer [5'.
TGTACCAATCTGAAGCGGAGCGCGCCCAATTTTTTTTTTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pTT3 vector. Library
went through one round of normalization to Cot5, and was
constructed by Bento Soares and M. Fatima Bonaldo."
ORIGIN
Query Match 80.9%; Score 17.8; DB 1; Length 271;
Best Local Similarity 90.5%; Pred. No. 1.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 2 GCCCTCTTCAGTCCAATCAAG 22
Db 251 GCTGCTTCAGTCCAATCAAG 231
RESULT 12
AA467041
LOCUS AA467041.1 NCI CGAP Kid12 Homo sapiens cDNA clone IMAGE:2873092
DEFINITION similar to TR:Q92478 Q92478 C-TYPE LECTIN. ;, mRNA sequence.
ACCESSION AA467041
VERSION AA467041.1 GI:7037147
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 289)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Akimura, T., Arakawa, T., Carninci, P., Furuno, M., Hanagaki, T.,
Hayatsu, N., Hiramoto, K., Hirooka, T., Hirozane, T., Imotani, K.,
Iehi, Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M.,
Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Numasaki, R.,
Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sakazume, N.,
Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,
Tanaka, T., Tomaru, A., Toya, T., Watahiki, A., Yasunishi, A.,
Muramatsu, M. and Hayashizaki, Y.

```

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 271)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbe@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: www-bio.lnl.gov/bbrp/image/image.html

Trace considered overall poor quality
Seq primer: -40UP from Gibco
High quality sequence stop: 1.
Location/Qualifiers

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1..271
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2873092"
/tissue_type="2 pooled tumors (clear cell type)"
/lab_host="DH10B"
/clone_lib="NCI-CGAP Kid12"
/note="Organ: kidney; Vector: pTT3D-Pac (Pharmacia) with
a modified polylinker; Site 1: Not I; Site 2: Eco RI;
Plasmid DNA from the normalized library NCI CGAP Kid5 was
prepared, and ss circles were made in vitro. Following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(cloneIDs 1323912-1325831, 1471368-1472903 and
1492104-1493255). Subtraction by Bento Soares and M.
Fatima Bonaldo."
ORIGIN
Query Match 80.9%; Score 17.8; DB 2; Length 271;
Best Local Similarity 90.5%; Pred. No. 1.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 2 GCCCTCTTCAGTCCAATCAAG 22
Db 153 GCCATCTTCAGTCCAATCCAG 173

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RESULT 13
BB720494
LOCUS BB720494.1 RIKEN full-length enriched, adult male liver tumor Mus
DEFINITION musculus cDNA clone C730036G02 3', mRNA sequence.
ACCESSION BB720494
VERSION BB720494.1 GI:16102067
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

```

```

REFERENCE 1 (bases 1 to 289)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Akimura, T., Arakawa, T., Carninci, P., Furuno, M., Hanagaki, T.,
Hayatsu, N., Hiramoto, K., Hirooka, T., Hirozane, T., Imotani, K.,
Iehi, Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M.,
Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Numasaki, R.,
Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sakazume, N.,
Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,
Tanaka, T., Tomaru, A., Toya, T., Watahiki, A., Yasunishi, A.,
Muramatsu, M. and Hayashizaki, Y.

```

TITLE RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al.
2001)
JOURNAL Unpublished (2001)
COMMENT Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.jp, URL: http://genome.gsc.riken.jp/
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M., and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. *Genome Res.* 10 (10), 1617-1630 (2000)
wagi, K., Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T.,
Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A.,
and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multiplexed sequencer. *Genome Res.*
10 (11), 1757-1771 (2000)
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P.,
Sugahara, Y., and Hayashizaki, Y.
Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)
Please visit our web site (<http://genome.gsc.riken.go.jp>) for
further details.
e mouse tissues.

FEATURES
source
Location/Qualifiers
1. .289
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="C730036G02"
/sex="male"
/issue_type="liver tumor"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="RIKEN full-length enriched, adult male liver
tumor"
/note="Site 1: SalI; Site 2: BamHI; cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
primed with a primer [5'
GAGAGAGAGCGCGCGCACTCGAGTTTCTTTTCTTTT 3'], cDNA was
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. Second strand cDNA was prepared with the
primer adapter of sequence [5'
GAGAGAGAGATTCGAGTTTCTTTTCTTTTATATCCCGCCCCCCC 3']. cDNA
was cleaved with BamHI and XhoI. Vector: a modified
pBluescript KS(+) after bulk excision from Lambda FLC I.
Tissue was provided by William A. Held, Roswell Park
Cancer Institute, Department of Molecular and Cellular
Biology, Elm and Carlton Streets, Buffalo, NY 14263, whose
assistance we gratefully acknowledge."

ORIGIN
Query Match 80.9%; Score 17.8; DB 2; Length 289;
Best Local Similarity 90.5%; Pred. No. 1.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAGTCCCAATCAA 21
|||||
Db 10 AGCCCTCTTCATTCCACTCAA 30
|||||

RESULT 14
BM151302
LOCUS
DEFINITION
TCBAP1D6134 Pediatric pre-B cell acute lymphoblastic leukemia
Baylor-HGSC project=TCBA Homo sapiens cDNA clone TCBAP6134, mRNA
sequence.
BM151302
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 295)
REFERENCE
AUTHORS
Wei, Y., Tsang, Y.T.M., Mei, G., Ku, J.M., Ali-Osman, F.R. Jr.,
Günatratne, P.H., Muzny, D., Bouck, J., Gibbs, R.A. and Margolin, J.F.
Pediatric Leukemia cDNA Sequencing Project (2001)
Unpublished (2001)
TITLE
JOURNAL
COMMENT
Contact: Dr. Judith F. Margolin
Texas Children's Cancer Center and Human Genome Sequencing Center
at Baylor College of Medicine
1102 Bates, MC3-3320 Houston, TX 77030, USA
Tel: 832-824-4536
Fax: 832-825-4038
Email: clones@ccc.org
Seq primer: M13 primer.
Location/Qualifiers
1. .295
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="TCBAP6134"
/sex="male"
/issue_type="leukopheresis"
/cell_type="pre-B cell"
/dev_stage="pediatric 2 years"
/lab_host="DH10B"
/clone_lib="Pediatric pre-B cell acute lymphoblastic
leukemia Baylor-HGSC project=TCBA"
/note="Vector: lambda pSB; Site 1: BamHI; Site 2: EcoRI;
First strand cDNA was primed with an anchored
XhoI-oligo(dT) primer [5'GGAGACTCGCGCGCGAGGAGGAG(T)VN
3'; V=A,C,G; N=A,C,G,T] and then dg tailed. Second strand
was primed with a BamHI-dc primer
[5'AGAGAGCTCGATCCGCGCGCAATATATATATAT(C) 3'].
Double-stranded cDNA was then digested with BamHI and XhoI
and directionally cloned into the BamHI and SalI sites of
lambda pSB vector. Library was constructed by Mei Yu at RIKEN
normalization. Library was constructed by Mei Yu at RIKEN
of Japan (Carninci P, Westover A, Nishiyama Y, Ohsumi T,
Itoh M, Nagaoka S, Sasaki, Okazaki Y, Muramatsu M,
Schneider C, Hayashizaki Y, High efficiency selection of
full-length cDNA by improved biotinylated cap trapper.,
DNA Res 4: 1, 61-6, Feb 28, 1997)"

ORIGIN
Query Match 80.9%; Score 17.8; DB 4; Length 295;
Best Local Similarity 90.5%; Pred. No. 1.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 2 GCCCTCTTCAGTCCCAATCAA 22
|||||
Db 269 GCCATCTTCAGTCCCAATCCAG 289
|||||

RESULT 15
AI203209/c
LOCUS
DEFINITION
q23h08.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:1941759 3',
mRNA sequence.
AI203209
ACCESSION
VERSION
KEYWORDS
EST.

SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 309)
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
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 Seq primer: -40UP from Gibco
 High quality sequence stop: 295.
FEATURES Location/Qualifiers
 1..309
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1941759"
 /tissue_type="pooled germ cell tumors"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP GC6"
 /note="Vector: pTT3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not 1; Site_2: Eco RI; Plasmid DNA from the normalized library NCI CGAP GC4 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIDs 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by Bento Soares and M. Fatima Bonaldo."
ORIGIN
 Query Match 80.9%; Score 17.8; DB 1; Length 309;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 GCCCTCTTCAGTCCAATCAAG 22
 ||| |||||
 Db 244 GCTGCTTCAGTCCAATCAAG 224
 T46996
 LOCUS T46996 310 bp mRNA linear EST 01-FEB-1995
 DEFINITION Yb12b06.s1 Stratagene placenta (#937225) Homo sapiens cDNA clone IMAGE:70931 3' similar to similar to SP:CD69_HUMAN Q07108 EARLY ACTIVATION ANTIGEN CD69, mRNA sequence.
 ACCESSION T46996
 VERSION T46996.1 GI:648979
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 310)
 Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiapelli, B., Chisoe, S., Dietrich, N., Dubuque, T., Favello, A., Gish, W., Hawkins, M., Hultman, M., Kucaba, T., Lacy, M., Le, M., Le, N., Mardis, E., Moore, B., Morris, M., Parsons, J., Prange, C., Rifkin, L., Rohlfing, T., Schellenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J., Trevaskis, E., Underwood, K., Wohlmann, P., Waterston, R., Wilson, R. and Marra, M.

TITLE Generation and analysis of 280,000 human expressed sequence tags
JOURNAL Genome Res. 6 (9), 807-828 (1996)
MEDLINE 97044478
PUBMED 889549
COMMENT Other ESTs: yb12b06.r1
 Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 Insert Size: 477
 High quality sequence stops: 272 Source: IMAGE Consortium, LLNL This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Possible reversed clone: polyI not found
 Insert Length: 477 Std Error: 0.00
 Seq primer: -21m13
 High quality sequence stop: 272.
FEATURES Location/Qualifiers
 1..310
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 /mol_type="mRNA"
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 /db_xref="taxon:9606"
 /clone="IMAGE:70931"
 /sex="male"
 /lab_host="SOLR cells (kanamycin resistant)"
 /clone_lib="Stratagene placenta (#937225)"
 /note="Organ: placenta; Vector: p Bluescript SK-; Site_1: EcoRI; Site_2: XhoI; Cloned unidirectionally. Primer: Oligo dt. Caucasian. Average insert size: 1.2 kb; Uni-ZAP XR Vector: -5' adaptor sequence: 5' GAATTGGCAGCAG 3' -3' adaptor sequence: 5' CTCGAGTCTTTTCTTTT 3"
ORIGIN
 Query Match 80.9%; Score 17.8; DB 7; Length 310;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 GCCCTCTTCAGTCCAATCAAG 22
 ||| |||||
 Db 175 GCCATCTTCAGTCCAATCCAG 195
 RESULT 17
 AA969466
 LOCUS AA969466 328 bp mRNA linear EST 07-JUL-1998
 DEFINITION o081d01.s1 NCI CGAP Kid5 Homo sapiens cDNA clone IMAGE:1572577 3' similar to TR:Q92478 Q92478 C-TYPE LECTIN.; mRNA sequence.
 ACCESSION AA969466
 VERSION AA969466.1 GI:3144646
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 328)
REFERENCE NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
TITLE Unpublished (1997)
JOURNAL Contact: Robert Strausberg, Ph.D.
COMMENT Email: cgapbs-remail.nih.gov
 Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 DNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
 Insert Length: 1094 Std Error: 0.00
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 1.

FEATURES

Location/Qualifiers
 1..328
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clones="IMAGE:1572577"
 /tissue_type="2 pooled tumors (clear cell type)"
 /lab_host="DH10B"
 /clone_lib="NCI_CGAP_Kid5"
 /notes="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' AACTGGAAGATTCGGCGCGCAATATTTTCTTTTCTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 80.9%; Score 17.8; DB 1; Length 328;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GCCCTCTTCAGTCCCAATCAAG 22

Db 153 GCCATCTTCAGTCCCAATCCAG 173

RESULT 18

AW615159/c
 LOCUS hg73h02.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:2951283 3',
 DEFINITION mRNA sequence.
 ACCESSION AW615159.1 GI:7320345
 VERSION AW615159
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 328)
 REFERENCE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 TITLE Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael R.
 R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.
 DNA Sequencing by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
 image.llnl.gov/image/html/iresources.shtml
 Seq primer: -40UP from Gibco.

FEATURES

Location/Qualifiers
 1..328
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clones="IMAGE:2951283"
 /tissue_type="pooled germ cell tumors"
 /lab_host="DH10B"
 /clone_lib="NCI_CGAP_GC6"
 /notes="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA

from the normalized library NCI_CGAP_GC4 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIda 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by Bento Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 80.9%; Score 17.8; DB 2; Length 328;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GCCCTCTTCAGTCCCAATCAAG 22

Db 230 GCTGCTTCAGTCCCAATCAAG 210

RESULT 19

AI307235
 LOCUS t518c07.x1 NCI_CGAP Kid12 Homo sapiens cDNA clone IMAGE:2054700 3',
 DEFINITION similar to TR:Q92478 Q92478 C-TYPE LECTIN. ;, mRNA sequence.
 ACCESSION AI307235.1 GI:4001991
 VERSION AI307235
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 339)
 REFERENCE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 TITLE Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 DNA Sequencing by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
 www-bio.llnl.gov/bbrp/image/image.html
 Insert length: 603 Std Error: 0.00
 Seq primer: -40UP from Gibco.

FEATURES

Location/Qualifiers
 1..339
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clones="IMAGE:2054700"
 /tissue_type="2 pooled tumors (clear cell type)"
 /lab_host="DH10B"
 /clone_lib="NCI_CGAP Kid12"
 /notes="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI;
 Plasmid DNA from the normalized library NCI_CGAP Kid5 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIda 1323912-1325831, 1471368-1472903 and 1492104-1493255). Subtraction by Bento Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 80.9%; Score 17.8; DB 1; Length 339;
 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GCCCTCTTCAGTCCCAATCAAG 22

QY 2 GCCCTCTTCAGTCCCAATCAAG 22
 Db 244 GCTGTCTTCAGTCCCAATCAAG 224

RESULT 25
 T46995/c

LOCUS
 DEFINITION Yb12b06.r1 Stratagene placenta (#937225) Homo sapiens cDNA clone
 IMAGE:70931 5' similar to similar to SP:CD69_HUMAN Q07108 EARLY
 ACTIVATION ANTIGEN CD69, mRNA sequence.

ACCESSION T46995
 VERSION
 KEYWORDS
 SOURCE EST.
 ORGANISM Homo sapiens (human)

REFERENCE
 AUTHORS
 1 (bases 1 to 339)
 Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiapelli, B.,
 Chisoe, S., Dietrich, N., Dubuque, T., Favello, A., Gish, W.,
 Hawkins, M., Hultman, M., Kucaba, T., Lacy, M., Le, M., Le, N.,
 Mardis, E., Moore, B., Morris, M., Parsons, J., Prange, C., Rifkin, L.,
 Rohlfing, T., Schellenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J.,
 Trevasakis, E., Underwood, K., Wohldmann, P., Waterston, R., Wilson, R.
 and Marra, M.

TITLE Generation and analysis of 280,000 human expressed sequence tags
 JOURNAL Genome Res. 6 (9), 807-828 (1996)
 MEDLINE 97044478
 PUBMED 8889549

COMMENT
 Other ESTs: yb12b06.s1
 Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 Insert Size: 477
 High quality sequence stops: 243
 Source: IMAGE Consortium, LLNL This
 clone is available royalty-free through LLNL ; contact the IMAGE
 Consortium (info@image.llnl.gov) for further information.
 Insert Length: 477 Std Error: 0.00
 Seq primer: M13RP1
 High quality sequence stop: 243.

FEATURES
 source
 1..339
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="GDB:491828"
 /db_xref="taxon:9606"
 /clone="IMAGE:70931"
 /sex="male"
 /lab_host="SOLR cells (kanamycin resistant)"
 /clone_lib="Stratagene placenta (#937225)"
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 Best Local Similarity 90.5%; Pred. No. 1.2e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GCCCTCTTCAGTCCCAATCAAG 22
 Db 118 GCCATCTTCAGTCCCAATCCAG 98

Search completed: September 6, 2005, 21:55:54
 Job time : 1578.31 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 770.688 Seconds
(without alignments)
1383.200 Million cell updates/sec

Title: US-10-729-421-35
Perfect score: 22
Sequence: 1 agcccttcagtcacatcaag 22

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 100 summaries

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11: gb_sts.*
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13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
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| C 2 | 22 | 100.0 | 1648 | 14 | AF375044 West Nile |
| C 3 | 22 | 100.0 | 1648 | 14 | AF375223 West Nile |
| C 4 | 22 | 100.0 | 2440 | 14 | AF194117 West Nile |
| C 5 | 22 | 100.0 | 10945 | 14 | AF202541 West Nile |
| C 6 | 22 | 100.0 | 10975 | 14 | AF206518 West Nile |
| C 7 | 22 | 100.0 | 10989 | 14 | AF268133 West Nile |
| C 8 | 22 | 100.0 | 10998 | 14 | AY278441 West Nile |
| C 9 | 22 | 100.0 | 11029 | 6 | AX576542 Sequence |
| C 10 | 22 | 100.0 | 11029 | 6 | AX577796 Sequence |
| C 11 | 22 | 100.0 | 11029 | 14 | AB185914 West Nile |
| C 12 | 22 | 100.0 | 11029 | 14 | AB185915 West Nile |
| C 13 | 22 | 100.0 | 11029 | 14 | AB185916 West Nile |
| C 14 | 22 | 100.0 | 11029 | 14 | AB185917 West Nile |
| C 15 | 22 | 100.0 | 11029 | 14 | AF196835 West Nile |
| C 16 | 22 | 100.0 | 11029 | 14 | AF260967 West Nile |
| C 17 | 22 | 100.0 | 11029 | 14 | AF404753 West Nile |
| C 18 | 22 | 100.0 | 11029 | 14 | AF404754 West Nile |
| C 19 | 22 | 100.0 | 11029 | 14 | AF404755 West Nile |

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| C 21 | 22 | 100.0 | 11029 | 14 | AF481864 West Nile |
| C 22 | 22 | 100.0 | 11029 | 14 | AF533540 West Nile |
| C 23 | 22 | 100.0 | 11029 | 14 | AY289214 West Nile |
| C 24 | 22 | 100.0 | 11057 | 14 | AY688948 West Nile |
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| C 27 | 21 | 95.5 | 2323 | 14 | AF130362 West Nile |
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| C 34 | 21 | 95.5 | 11028 | 14 | AY490240 West Nile |
| C 35 | 21 | 95.5 | 11029 | 14 | AF260968 West Nile |
| C 36 | 21 | 95.5 | 11029 | 14 | AF260969 West Nile |
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| C 40 | 19.4 | 88.2 | 11022 | 14 | AY274505 |
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| C 53 | 18.4 | 83.6 | 238643 | 2 | AC133255 Rattus no |
| C 54 | 18.4 | 83.6 | 275209 | 2 | AC097164 Rattus no |
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| C 60 | 17.8 | 80.9 | 380 | 6 | BD119800 EST and e |
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ALIGNMENTS

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 ACCESSION AF375042
 VERSION AF375042.1 GI:19421847
 KEYWORDS
 SOURCE West Nile virus
 ORGANISM West Nile virus
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.
 REFERENCE 1 (bases 1 to 1648)
 AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Weiss,L., Grossman,Z. and Bin,H.
 TITLE Isolation and characterization of West Nile virus from the blood of viremic patients during the 2000 outbreak in Israel
 JOURNAL Emerging Infect. Dis. 7 (4), 748-750 (2001)
 MEDLINE 21469825
 PUBMED 11585544
 REFERENCE 2 (bases 1 to 1648)
 AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Weiss,L., Grossman,Z. and Bin,H.
 TITLE Direct Submission
 JOURNAL Submitted (30-APR-2001) Central Virology Laboratory, Ministry of Health, Public Health Laboratories, Sheba Medical Center, Tel Hashomer 52621, Israel

FEATURES

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 VERSION AF375044.1 GI:19421851
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 SOURCE West Nile virus
 ORGANISM West Nile virus
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.
 REFERENCE 1 (bases 1 to 1648)
 AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Weiss,L., Grossman,Z. and Bin,H.
 TITLE Isolation and characterization of West Nile virus from the blood of viremic patients during the 2000 outbreak in Israel
 JOURNAL Emerging Infect. Dis. 7 (4), 748-750 (2001)
 MEDLINE 21469825
 PUBMED 11585544
 REFERENCE 2 (bases 1 to 1648)
 AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Weiss,L., Grossman,Z. and Bin,H.
 TITLE Direct Submission
 JOURNAL Submitted (30-APR-2001) Central Virology Laboratory, Ministry of Health, Public Health Laboratories, Sheba Medical Center, Tel Hashomer 52621, Israel

FEATURES

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 Best Local Similarity 100.0%; Pred. No. 1.5;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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RESULT 3

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 ACCESSION AF375223
 VERSION AF375223.1 GI:17226060
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 SOURCE West Nile virus
 ORGANISM West Nile virus
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.
 REFERENCE 1 (bases 1 to 1648)
 AUTHORS Banet,C., Brill,A., Samina,I., Yadin,H., Straum,Y., Weisman,J., Pokamonski,S., King,R., Deubel,V. and Malkinson,M.
 TITLE Phylogenetic relationships of West Nile viruses isolated in Israel

REFERENCE

1 (bases 1 to 1648)
 Banet,C., Brill,A., Samina,I., Yadin,H., Straum,Y., Weisman,J., Pokamonski,S., King,R., Deubel,V. and Malkinson,M.
 Phylogenetic relationships of West Nile viruses isolated in Israel

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from 1997 to 2000
Unpublished
REFERENCE 2 (bases 1 to 1648)
AUTHORS Banet,C., Brill,A., Samina,I., Yadin,H., Straum,Y., Weisman,J.,
Pokamonski,S., King,R., Deubel,V. and Malkinson,M.
TITLE Direct Submission
JOURNAL Submitted (01-MAY-2001) Kimron Veterinary Institute, Beit Degan
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DEFINITION AF194117
ACCESSION AF194117.1 GI:6715269
VERSION
KEYWORDS
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ORGANISM
REFERENCE 1 (sites)
AUTHORS Lanciotti,R.S., Roehrig,J.T., Deubel,V., Smith,J., Parker,M.,
Steele,K., Crise,B., Volpe,K.E., Crabtree,M.B., Scherret,J.H.,
Hall,R.A., Mackenzie,J.S., Cropp,C.B., Panigrahy,B., Ostlund,E.,
Schmitt,B., Malkinson,M., Banet,C., Weisman,J., Komar,N.,
Savage,H.M., Stone,W., McNamara,T. and Gubler,D.J.
TITLE Origin of the West Nile virus responsible for an outbreak of
encephalitis in the northeastern United States
JOURNAL Science 286 (5448), 2333-2337 (1999)
MEDLINE 20070288
PUBMED 10600742
REFERENCE 2 (bases 1 to 2440)
AUTHORS Parker,M.D., Crise,B.J., Clayton,J.M. and Smith,J.F.
TITLE Direct Submission
JOURNAL Submitted (13-OCT-1999) Virology Division, U.S. Army Medical
Research Institute of Infectious Diseases, Bldg. 1425 Fort Detrick,
Frederick, Maryland 21702, USA
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ACCESSION AF202541
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 10945)
AUTHORS Jia,X.Y., Brice,T., Jordan,I., Rambaut,A., Chi,H.C.,
Mackenzie,J.S., Hall,R.A., Scherret,J. and Lipkin,W.I.
TITLE Genetic analysis of West Nile New York 1999 encephalitis virus
JOURNAL Lancet 354 (9194), 1971-1972 (1999)
MEDLINE 20086017
PUBMED 10622305
REFERENCE 2 (bases 1 to 10945)
AUTHORS Jia,X.Y., Brice,T., Jordan,I. and Lipkin,W.I.
TITLE Direct Submission
JOURNAL Submitted (06-NOV-1999) Emerging Diseases Laboratory, Dept.
Microbiology & Molecular Genetics and Neurology, University of
California, Irvine, 3101 Gillespie Neuroscience Facility, Irvine,
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1 (bases 1 to 10975)
Wakem, E.M., French, R.A., Garmendia, A.E. and Van Kruiningen, H.J.
Isolation of West Nile virus from mosquitoes, crows, and a Cooper's
hawk in Connecticut
Science 286 (5448), 2331-2333 (1999)
JOURNAL
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REFERENCE 2 (bases 1 to 10975)
Vossbrinck, C.R., Anderson, J.F. and Andreadis, T.G.
AUTHORS
TITLE Genome Sequence of West Nile Virus from Culex pipiens isolate
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 10975)
Anderson, J.F., Andreadis, T.G. and Vossbrinck, C.R.
AUTHORS
TITLE Direct Submission
JOURNAL Direct Submission
REFERENCE 4 (bases 1 to 10975)
Anderson, J.F., Andreadis, T.G. and Vossbrinck, C.R.
AUTHORS
TITLE Submitted (08-MAY-2000) Soil and Water, Connecticut Agricultural
Experiment Station, 123 Huntington Street, New Haven, CT 06511, USA
JOURNAL Experiment Station, 123 Huntington Street, New Haven, CT 06511, USA
REMARK Sequence update by submitter
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West Nile virus strain PAH001 polyprotein (pol) gene, complete cds.
AY268133
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West Nile virus (WNV)
West Nile virus
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1 (bases 1 to 10989)
Charrel,R.N., Brault,A.C., Gallian,P., Lemaason,J.-J., Murque,B.,
Murri,S., Pastorino,B., Zeller,H., de chessee,R., de Micco,P. and de
Lamballerie,X.
Evolutionary relationship between Old World West Nile virus
strains. Evidence for viral gene flow between africa, the middle
east, and europe
Virology 315 (2), 381-388 (2003)
22949215
14585341
2 (bases 1 to 10989)
de Lamballerie,X., Brault,A.C., Gallian,P., Lemaason,J., Murque,B.,
Murri,S., Pastorino,B., Zeller,H., Dechesse,R., de Micco,P. and
Charrel,R.N.
Direct Submission
Submitted (03-APR-2003) Virology, Medical University, 27 bd Jean
Moulin, Marseille 13005, France
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ORIGIN

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| Query Match | 100.0%; | Score 22; | DB 14; | Length 10989; |
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| DEFINITION | West Nile virus isolate Ast99-901, complete genome. | | | |
| ACCESSION | AY278441 | | | |
| VERSION | AY278441.1 | GI:30349729 | | |
| KEYWORDS | | | | |
| SOURCE | West Nile virus (WNV) | | | |
| ORGANISM | West Nile virus | | | |
| | Flavivirus; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; | | | |
| | Flavivirus; Japanese encephalitis virus group. | | | |
| REFERENCE | 1 | (bases 1 to 10998) | | |
| AUTHORS | Voronina, A.G., Prilipov, A.G., Kinney, R.M., Samokhvalov, E.I., | | | |
| | Savage, H.M., Alkhovsky, S.V., Tsychia, R., Sadykova, G.K., | | | |
| | Shatalov, A.G., Usachev, E.V., Mokhonov, V.V., Butenko, A.M., | | | |
| | Larichev, V.F., Gubler, D.J. and Lvov, D.K. | | | |
| | Analysis of a new variants of West Nile virus | | | |
| TITLE | Unpublished | | | |
| JOURNAL | | | | |
| REFERENCE | 2 | (bases 1 to 10998) | | |
| AUTHORS | Voronina, A.G., Prilipov, A.G., Kinney, R.M., Samokhvalov, E.I., | | | |
| | Savage, H.M., Alkhovsky, S.V., Tsychia, R., Sadykova, G.K., | | | |
| | Shatalov, A.G., Usachev, E.V., Mokhonov, V.V., Butenko, A.M., | | | |
| | Larichev, V.F., Gubler, D.J. and Lvov, D.K. | | | |
| | Direct Submission | | | |
| TITLE | | | | |
| JOURNAL | | | | |
| FEATURES | Submitted (17-APR-2003) Molecular Genetic, Ivanovsky Virology | | | |
| | Institute, Gamalei 16, Moscow 123098, Russia | | | |
| | Location/Qualifiers | | | |
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CDS

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Query Match 100.0%; Score 22; DB 14; Length 10998;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
    |||||
Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 9
AX576542/c
LOCUS AX576542 11029 bp DNA linear PAT 08-JAN-2003
DEFINITION Sequence 1 from Patent WO02081511.
ACCESSION AX576542
VERSION AX576542.1 GI:27646162
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS Despres, P., Deubel, V., Guenet, J.L., Drouet, M.T., Malkinson, M.K.,
Banet, C.K., Frenkel, M.P., Courageot, M.P., Coulibaly, F.,
Catteau, A., Flanand, M., Weber, P., and Ceccaldi, P.E.,
Neurovirulent strain of the west nile virus and applications
thereof
JOURNAL Patent: WO 02081511-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR) ; Kimron Veterinary Institute (IL)
FEATURES
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1. .11029
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Best Local Similarity 100.0%; Pred. No. 1.5;
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Db 195 AGCCCTCTTCAGTCCAATCAAG 174

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DEFINITION Sequence 1 from Patent WO02081741.
ACCESSION AX577796
VERSION AX577796.1 GI:27647035
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS Guenet, J.L., Mashimo, T., Simon-Chazottes, D., Montagutelli, X.,
Frenkel, M.P., Despres, P., Deubel, V., Bonhomme, F., and Lucas, M.
Use of products of genes of the 2'-5' oligoadenylate synthetase
family (oas) for screening antiviral agents and for detecting
responsiveness to flaviviridae infection
Patent: WO 02081741-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR) ; CENTRE NATIONAL DE LA RECHERCHE
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ORIGIN

Query Match 100.0%; Score 22; DB 6; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 1.5; Indels 0; Gaps 0;
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 VERSION AB185914.2 GI:50872124
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 SOURCE West Nile virus (WNV)

ORGANISM West Nile virus
 REFERENCE 1 Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae; Flavivirus; Japanese encephalitis virus group.
 AUTHORS Shirato, K., Miyoshi, H., Goto, A., Ako, Y., Ueki, T., Kariwa, H. and Takashima, I.
 TITLE Correlation between viral envelope glycosylation and neuroinvasiveness of the New York strain of the West Nile virus
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 11029)
 AUTHORS Shirato, K., Kariwa, H. and Takashima, I.
 TITLE Direct Submission
 JOURNAL Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of Veterinary Medicine, Hokkaido University, Laboratory of Public Health, Department of Environmental Veterinary Medicine; Kita-19 Nishi-5, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
 COMMENT (E-mail: shirato@vetmed.hokudai.ac.jp. Tel: 81-11-706-5213), Fax: 81-11-706-5213
 ON Jul 30, 2004 this sequence version replaced gi:50838778.
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 Location/Qualifiers
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 1.5;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
 |||||
 Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 12

AB185915/c

LOCUS

DEFINITION

West Nile virus gene for polyprotein precursor protein, complete cds, isolate: 6-SP.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

West Nile virus (WNV)
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.

REFERENCE

AUTHORS

Shirato, K., Miyoshi, H., Goto, A., Ako, Y., Ueki, T., Kariwa, H. and Takashima, I.

TITLE

Correlation between viral envelope glycosylation and neuroinvasiveness of the New York strain of the West Nile virus

JOURNAL

AUTHORS

TITLE

JOURNAL

Unpublished
 2 (bases 1 to 11029)
 Shirato, K., Kariwa, H. and Takashima, I.
 Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of Veterinary Medicine, Hokkaido University, Laboratory of Public Health, Department of Environmental Veterinary Medicine, Kita-19 Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
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COMMENT

FEATURES

source

On Jul 30, 2004 this sequence version replaced gi:50838780.

Location/Qualifiers

1. 11029

/organism="West Nile virus"

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CDS

ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 1.5;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22

|||||

Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 13

AB185916/c

LOCUS

DEFINITION

West Nile virus gene for polyprotein precursor protein, complete cds, isolate: B-SP.

ACCESSION

AB185916

VRL 30-JUL-2004

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VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source
CDS

AB185916.1 GI:50838782
West Nile virus (WNV)
West Nile virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1
Shirato,K., Miyoshi,H., Goto,A., Ako,Y., Ueki,T., Kariwa,H. and
Takashima,I.
Correlation between viral envelope glycosylation and
neuroinvasiveness of the New York strain of the West Nile virus
unpublished
2 (bases 1 to 11029)
Shirato,K., Kariwa,H. and Takashima,I.
Direct Submission
Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of
Veterinary Medicine, Hokkaido University, Laboratory of Public
Health, Department of Environmental Veterinary Medicine; Kita-19
Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
(E-mail:shirato@vetmed.hokudai.ac.jp, Tel:81-11-706-5213),
Fax:81-11-706-5213)
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22

|||||

Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 14

AB185917/c

LOCUS

DEFINITION

West Nile virus gene for polyprotein precursor protein, complete

AB185917

AB185917.1 GI:50838784

West Nile virus (WNV)

West Nile virus

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

Flavivirus; Japanese encephalitis virus group.

1

Shirato,K., Miyoshi,H., Goto,A., Ako,Y., Ueki,T., Kariwa,H. and

Takashima,I.

Correlation between viral envelope glycosylation and

neuroinvasiveness of the New York strain of the West Nile virus

unpublished

2 (bases 1 to 11029)

Shirato,K., Kariwa,H. and Takashima,I.

Direct Submission

Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of

Veterinary Medicine, Hokkaido University, Laboratory of Public

Health, Department of Environmental Veterinary Medicine; Kita-19

Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan

(E-mail:shirato@vetmed.hokudai.ac.jp, Tel:81-11-706-5213),

Fax:81-11-706-5213)

Location/Qualifiers

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/organism="West Nile virus"

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97. .10398

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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
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Db 195 AGCCCTCTTCAGTCCAATCAAG 174
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RESULT 15
AF196835/c
LOCUS
DEFINITION West Nile virus strain NY99-flamingo382-99, complete genome.
ACCESSION AF196835

VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
REFERENCE
AUTHORS
TITLE
JOURNAL
REMARK
COMMENT
FEATURES
source

AF196835.2 GI:11597239
West Nile virus
West Nile virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 11029)
Lancioti,R.S., Roehrig,J.T., Deubel,V., Smith,J., Parker,M.,
Steel,K., Crise,B., Volpe,K.E., Crabtree,M.B., Scherret,J.H.,
Hall,R.A., Mackenzie,J.S., Cropp,C.B., Panigrahy,B., Ostlund,E.,
Schmitt,B., Malkinson,M., Banet,C., Weissman,J., Komar,N.,
Savage,H.M., Stone,W., McNamara,T. and Gubler,D.J.
Origin of the West Nile virus responsible for an outbreak of
encephalitis in the northeastern United States
Science 286 (5448), 2333-2337 (1999)
20070288
10600742
2 (bases 1 to 11029)
Lancioti,R., Roehrig,J., Volpe,K. and Panigrahy,B.
Direct Submission
Submitted (20-OCT-1999) Division of Vector-Borne Diseases, Centers
for Disease Control and Prevention, Rampart Road, Fort Collins, CO
80521, USA
3 (bases 1 to 11029)
Lancioti,R., Roehrig,J., Volpe,K. and Panigrahy,B.
Direct Submission
Submitted (07-DEC-2000) Division of Vector-Borne Diseases, Centers
for Disease Control and Prevention, Rampart Road, Fort Collins, CO.
80521, USA
Sequence update by submitter
On Dec 7, 2000 this sequence version replaced gi:6636174.
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 195 AGCCCTCTTCAGTCCAATCAAG 174
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RESULT 16
AF260967/c
LOCUS AF260967 11029 bp RNA linear VRL 27-AUG-2000
DEFINITION West Nile virus strain NY99-eghs, complete genome.
ACCESSION AF260967
VERSION AF260967.1 GI:9930133
KEYWORDS West Nile virus
SOURCE West Nile virus
ORGANISM West Nile virus

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 11029)
Bowen.M., Meyer.R.F., McKinney.N., Morrill.W. and Lanciotti.R.
Complete genomic sequence of West Nile virus equine isolate New
York 1999
Unpublished
2 (bases 1 to 11029)
Bowen.M., Meyer.R.F., McKinney.N., Morrill.W. and Lanciotti.R.
Direct Submission
Submitted (27-APR-2000) Arbovirus Diseases Branch, Centers for
Disease Control & Prevention, Rampart Road, Fort Collins, CO 80521,
USA
Location/Qualifiers
1. 11029
/organism="West Nile virus"
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTTTCAGTCCCAATCAAG 22
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Db 195 AGCCCTTTCAGTCCCAATCAAG 174
|||||

RESULT 17
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LOCUS West Nile virus isolate WN MD 2000-crow265, complete genome.
DEFINITION
ACCESSION AF404753
VERSION AF404753.1 GI:21929232
KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus
LORPATWISLYAVTAVTLTLLKHLITSYINTSLTSINVSALPTLARGPFPDV
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 11029)
Lanciotti,R.S., Ebel,G.D., Deubel,V., Kerst,A.J., Murri,S.,
Meyer,R., Bowen,M., McKinney,N., Morrill,W.E., Crabtree,M.B.,
Kramer,L.D. and Roehrig,J.T.
Complete genome sequences and phylogenetic analysis of West Nile
virus strains isolated from the United States, Europe, and the
Middle East
Virology 298 (1), 96-105 (2002)
JOURNAL MEDLINE 22089180
PUBMED 12093177
REFERENCE 2 (bases 1 to 11029)
Lanciotti,R.S., Ebel,G.D. and Kerst,A.J.
AUTHORS
TITLE Direct Submission
JOURNAL Submitted (02-AUG-2001) Division of Vector-Borne Infectious
Diseases, Centers for Disease Control & Prevention, Rampart Road,

FEATURES
source

Fort Collins, CO 80521, USA

Location/Qualifiers

1. 11029

/organism="West Nile virus"

/mol_type="genomic RNA"

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97.10398

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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
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Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 18

AF404754/c 11029 bp ss-RNA linear VRL 23-JUL-2002
LOCUS West Nile virus isolate WN NJ 2000 MQ5488, complete genome.

DEFINITION

AF404754

ACCESSION

AF404754.1

VERSION

1

KEYWORDS

West Nile virus

SOURCE

West Nile virus

ORGANISM

Flavivirus; Japanese encephalitis virus group.

REFERENCE

1 (bases 1 to 11029)

AUTHORS

Lanciotti, R.S., Ebel, G.D., Deubel, V., Kerst, A.J., Murri, S.,
Meyer, R., Bowen, M., McKinney, N., Morrill, W.E., Crabtree, M.B.,
Kramer, L.D. and Roehrig, J.T.

TITLE

Complete genome sequences and phylogenetic analysis of West Nile
virus strains isolated from the United States, Europe, and the
Middle East

JOURNAL

Virology 298 (1), 96-105 (2002)

MEDLINE

22089180

PUBMED

12093177

REFERENCE

2 (bases 1 to 11029)

AUTHORS

Lanciotti, R.S., Ebel, G.D. and Kerst, A.J.

TITLE

Direct Submission

JOURNAL

Submitted (02-AUG-2001) Division of Vector-Borne Infectious
Diseases, Centers for Disease Control & Prevention, Rampart Road,
Fort Collins, CO 80521, USA

LOCATION/Qualifiers

1..11029

FEATURES

Location/Qualifiers

1..11029

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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22

|||||

Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 19

AF404755/c

LOCUS

West Nile virus isolate WN NY 2000-grouse3282, complete genome.

DEFINITION

AF404755

ACCESSION

AF404755.1

VERSION

1

KEYWORDS

West Nile virus

SOURCE

West Nile virus

ORGANISM

Flavivirus; Japanese encephalitis virus group.

REFERENCE

1 (bases 1 to 11029)

AUTHORS

Lanciotti, R.S., Ebel, G.D., Deubel, V., Kerst, A.J., Murri, S.,
Meyer, R., Bowen, M., McKinney, N., Morrill, W.E., Crabtree, M.B.,
Kramer, L.D. and Roehrig, J.T.

TITLE

Complete genome sequences and phylogenetic analysis of West Nile
virus strains isolated from the United States, Europe, and the
Middle East

JOURNAL

Virology 298 (1), 96-105 (2002)


```

MEDLINE
PUBMED
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source

22089180
12093177
2 (bases 1 to 11029)
Ebel,G.D., Kerst,A.J. and Lanciotti,R.S.
Direct Submission
Submitted (02-AUG-2001) Division of Vector-Borne Infectious
Diseases, Centers for Disease Control & Prevention, Rampart Road,
Fort Collins, CO 80521, USA
Location/Qualifiers
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ORIGIN
Query Match 100.0%; Score 22; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAATCAAG 22
Db 195 AGCCCTCTTCAATCAAG 174
RESULT 20
AF404756/C
LOCUS AF404756 11029 bp ss-RNA linear VRL 23-JUL-2002
DEFINITION West Nile virus isolate WN NY 2000-crow3356, complete genome.
ACCESSION AF404756
VERSION AF404756.1 GI:21929238
KEYWORDS
SOURCE
ORGANISM
West Nile virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE
1 (bases 1 to 11029)
Lanciotti,R.S., Ebel,G.D., Deubel,V., Kerst,A.J., Murri,S.,
Meyer,R., Bowen,M., McKinney,N., Morrill,W.E., Crabtree,M.B.,
Kramer,L.D. and Roehrig,J.F.
Complete genome sequences and phylogenetic analysis of West Nile
virus strains isolated from the United States, Europe, and the
Middle East
Virology 298 (1), 96-105 (2002)
JOURNAL
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Ebel,G.D., Kerst,A.J. and Lanciotti,R.S.
Direct Submission
Submitted (02-AUG-2001) Division of Vector-Borne Infectious
Diseases, Centers for Disease Control & Prevention, Rampart Road,
Fort Collins, CO 80521, USA
Location/Qualifiers
1. .11029
/organism="West Nile virus"
/mol_type="genomic RNA"
/isolate="WN NY 2000-crow3356"
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97. .10398
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NS2b, NS3, NS4a, NS4b, and NS5"
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ORIGIN

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VERSION AF481864.1 GI:19387527
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ORGANISM West Nile virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE 1 (bases 1 to 11029)

AUTHORS Malkinson,M., Banet,C., Weisman,Y., Pokamunski,S., King,R.,
Drouet,M.T. and Deubel,V.
TITLE Introduction of West Nile virus in the Middle East by migrating
white storks
JOURNAL Emerging Infect. Dis. 8 (4), 392-397 (2002)
MEDLINE 21968420
PUBMED 11971773
REFERENCE 2 (bases 1 to 11029)
AUTHORS Deubel,V., Malkinson,M. and Banet,C.
TITLE Direct Submission
JOURNAL Submitted (08-FEB-2002) CERVI, Institut Pasteur, 21 Avenue Tony
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ORIGIN

Query Match 100.0%; Score 22; DB 14; Length 11029;

Best Local Similarity 100.0%; Pred. No. 1.5;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCAGTCCAATCAAG 22

Db 195 AGCCCTCTTCAGTCCAATCAAG 174

RESULT 22

AF533540/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

FEATURES

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West Nile virus strain TYP 8533 complete genome.
AY289214 GI:33948906
West Nile virus (WNV)
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 11029)
Granwehr, B.P., Li, L., Davis, C.T., Beasley, D.W.C. and Barrett, A.D.T.
Phylogenetic Analysis of a Human Isolate of West Nile Virus from
the Gulf Coast of Texas, 2002
Unpublished
2 (bases 1 to 11029)
Granwehr, B.P. and Barrett, A.D.T.
Direct Submission
Submitted (01-MAY-2003) Internal Medicine-Infectious Diseases,
University of Texas Medical Branch, 301 University Blvd., Rte.
0435, Galveston, TX 77555/0435, USA
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3' UTR
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Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCCCTCTTCACTCCAATCAAG 22
Db 195 AGCCCTCTTCACTCCAATCAAG 174

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LOCUS      AY688948 11057 bp      RNA      linear      VRL 15-AUG-2004
DEFINITION West Nile virus strain Sarafend, complete genome.
ACCESSION  AY688948
VERSION     AY688948.1  GI:51095221
KEYWORDS
SOURCE      West Nile virus (WNV)
ORGANISM    West Nile virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Flavivirus; Japanese encephalitis virus group.
REFERENCE   1 (bases 1 to 11057)
AUTHORS     Li,J., Bhuvanankantham,R. and Ng,M.-L.
TITLE       Construction and characterization of an infectious West Nile
            (Sarafend) clone
JOURNAL     Unpublished
REFERENCE   2 (bases 1 to 11057)
AUTHORS     Li,J., Bhuvanankantham,R. and Ng,M.-L.
TITLE       Direct Submission
JOURNAL     Submitted (18-JUL-2004) Microbiology, National University of
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Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCCCTCTTCAGTCCAATCAAG 22
Db 195 AGCCCTCTTCAGTCCAATCAAG 174
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LOCUS      AF375043 1648 bp      mRNA      linear      VRL 14-MAR-2002
DEFINITION West Nile virus isolate MN_0233 polyprotein mRNA, partial cds.
ACCESSION  AF375043
VERSION     AF375043.1  GI:19421849
KEYWORDS
SOURCE      West Nile virus
ORGANISM    West Nile virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Flavivirus; Japanese encephalitis virus group.
REFERENCE   1 (bases 1 to 1648)
AUTHORS     HindiyeH,M., Shulman,L.M., Mendelson,E., Weiss,L., Grosseman,Z. and
            Bin,H.
TITLE       Isolation and characterization of West Nile virus from the blood of
            viremic patients during the 2000 outbreak in Israel
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JOURNAL Emerging Infect. Dis. 7 (4), 748-750 (2001)
MEDLINE 21469825
PUBMED 11595544
REFERENCE 2 (bases 1 to 1648)
AUTHORS Hindiyyeh,M., Shulman,L.M., Mendelson,E., Grossman,Z., Weiss,L. and Bin,H.
TITLE Direct Submission
JOURNAL Submitted (30-APR-2001) Central Virology Laboratory, Ministry of Health, Public Health Laboratories, Sheba Medical Center, Tel Hashomer 52621, Israel
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Qy 2 GCCCTCTTCAGTCCCAATCAAG 22
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Search completed: September 6, 2005, 20:29:45
Job time : 775.688 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 735.656 Seconds
(without alignments)
1383.200 Million cell updates/sec

Title: US-10-729-421-34
Perfect score: 21
Sequence: 1 ccgggtgtcaatgctaaa 21

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

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3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_ro.*

11: gb_ats.*

12: gb_ay.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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| 2 | 21 | 100.0 | 2440 | 14 AF194117 | AF194117 West Nile |
| 3 | 21 | 100.0 | 10664 | 14 KUNCG | D00246 Kunjin viru |
| 4 | 21 | 100.0 | 10842 | 14 AY278442 | AY278442 West Nile |
| 5 | 21 | 100.0 | 10845 | 14 AY27252 | AY27252 West Nile |
| 6 | 21 | 100.0 | 10945 | 14 AF202541 | AF202541 West Nile |
| 7 | 21 | 100.0 | 10962 | 14 WNFCC | M12294 West Nile v |
| 8 | 21 | 100.0 | 10972 | 14 AF317203 | AF317203 West Nile |
| 9 | 21 | 100.0 | 10975 | 14 AF206518 | AF206518 West Nile |
| 10 | 21 | 100.0 | 10984 | 14 AY262283 | AY262283 West Nile |
| 11 | 21 | 100.0 | 10989 | 14 AY268132 | AY268132 West Nile |
| 12 | 21 | 100.0 | 10989 | 14 AY268133 | AY268133 West Nile |
| 13 | 21 | 100.0 | 11022 | 14 AY274504 | AY274504 Kunjin vi |
| 14 | 21 | 100.0 | 11022 | 14 AY274505 | AY274505 Kunjin vi |
| 15 | 21 | 100.0 | 11028 | 14 AY490240 | AY490240 West Nile |
| 16 | 21 | 100.0 | 11029 | 6 AY576542 | AY576542 Sequence |
| 17 | 21 | 100.0 | 11029 | 6 AY577796 | AY577796 Sequence |
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| 84 | 17.8 | 84.8 | 10976 | 14 JEVBEICG | L48961 Japanese en |
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| 86 | 17.8 | 84.8 | 10976 | 14 JEVSAV | D90195 Japanese en |
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| 91 | 17.8 | 84.8 | 11014 | 14 AF161266 | AF161266 Murray Va |
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 C 95 17.4 82.9 150121 10 AC114925
 C 96 17.4 82.9 197486 2 AC145692
 C 97 17.4 82.9 199087 2 CR848021
 C 98 17.4 82.9 200081 2 AC107693
 C 99 17.4 82.9 200890 5 BX465192
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ALIGNMENTS

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 DEFINITION M32560
 ACCESSION M32560.1 GI:336165

KEYWORDS
 SOURCE West Nile virus
 ORGANISM West Nile virus
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 Flavivirus; Japanese encephalitis virus group.

REFERENCE
 AUTHORS Castle, E. and Wengler, G.

TITLE Nucleotide sequence of the 5'-terminal untranslated part of the

JOURNAL genome of the flavivirus West Nile virus

MEDLINE Arch. Virol. 922, 309-313 (1987)

COMMENT Original source text: West Nile virus cDNA to genomic RNA.

FEATURES
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LOCUS West Nile virus structural protein precursor, gene, partial cds.

DEFINITION AF194117

ACCESSION AF194117.1 GI:6715269

VERSION

KEYWORDS

SOURCE West Nile virus

ORGANISM West Nile virus

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

Flavivirus; Japanese encephalitis virus group.

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

REFERENCE

AUTHORS

TITLE

JOURNAL

FEATURES

source

CDS

ORIGIN

Query Match

Best Local Similarity

Matches

Qy

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RESULT 3

KUNCG

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

1 (sites)

Lancioti, R.S., Roehrig, J.T., Deubel, V., Smith, J., Parker, M.,

Steele, K., Crise, B., Volpe, K.E., Crabtree, M.B., Scherret, J.H.,

Hall, R.A., Mackenzie, J.S., Cropp, C.B., Panigrahy, B., Ostlund, E.,

Schmitt, B., Malkinson, M., Banet, C., Weissman, J., Komar, N.,

Savage, H.M., Stone, W., McNamara, T., and Gubler, D.J.

Origin of the West Nile virus responsible for an outbreak of

encephalitis in the northeastern United States

Science 286 (5448), 2333-2337 (1999)

20070288

10600742

2 (bases 1 to 2440)

Parker, M.D., Crise, B.J., Clayton, J.M. and Smith, J.F.

Direct Submission

Submitted (13-OCT-1999) Virology Division, U.S. Army Medical

Research Institute of Infectious Diseases, Bldg. 1425 Fort Detrick,

Frederick, Maryland 21702, USA

Location/Qualifiers

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100.0%; Score 21; DB 14; Length 2440;

Best Local Similarity 100.0%; Pred. No. 1.4;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21

Db 33 CCGGGCTGTCATATGCTAAA 53

KUNCG

Kunjin virus gene for polyprotein (C, prM, E, NS1, NS2A, NS2B, NS3,

NS4A, NS4B, NS5), complete cds.

D00246

D00246.1 GI:221966

M (membrane protein); prM (precursor of M); NS5; NS4B; NS4A; NS3;

NS2B; NS2A; NS1; E (envelope protein); C (core protein);

polyprotein.

Kunjin virus

Kunjin virus

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

Flavivirus; Japanese encephalitis virus group.

1 (bases 1 to 10664)

Coia, G., Parker, M.D., Speight, G., Byrne, M.E. and Westaway, E.G.

Nucleotide and complete amino acid sequences of Kunjin virus:

definitive gene order and characteristics of the virus-specified

proteins

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Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CCGGGCTGTCAATATGCTAAA 21
Db 129 CCGGGCTGTCAATATGCTAAA 149
RESULT 5
LOCUS AY277252 10845 bp RNA linear VRL 03-MAY-2003
DEFINITION West Nile virus isolate LEIV-Vlg99-27889, complete genome.
ACCESSION AY277252
VERSION AY277252.1 GI:30349727
KEYWORDS
SOURCE West Nile virus (WNV)
ORGANISM West Nile virus
REFERENCE 1 (bases 1 to 10845)
AUTHORS Prilipov,A.G., Kinney,R.M., Samokhvalov,E.I., Savage,H.M.,
Alkhovsky,S.V., Tsychia,R., Gromashevsky,V.L., Sadykova,G.K.,
Shatalov,A.G., Usachev,E.V., Mokhonov,V.V., Voronina,A.G.,
Butenko,A.M., Larichev,V.F., Gubler,D.J. and Lvov,D.K.
TITLE Analysis of a new variants of West Nile virus
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 10845)
AUTHORS Prilipov,A.G., Kinney,R.M., Samokhvalov,E.I., Savage,H.M.,
Alkhovsky,S.V., Tsychia,R., Gromashevsky,V.L., Sadykova,G.K.,
Shatalov,A.G., Usachev,E.V., Mokhonov,V.V., Voronina,A.G.,
Butenko,A.M., Larichev,V.F., Gubler,D.J. and Lvov,D.K.
TITLE Direct Submision
JOURNAL Submitted (15-APR-2003) Molecular Genetic, Ivanovsky Virology
Institute, Gamalei 16, Moscow 123098, Russia
FEATURES
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ORIGIN
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Best Local Similarity 100.08; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGCTCAATATGCTAAA 21
Db 129 CCGGGCTGCTCAATATGCTAAA 149
RESULT 8
AF317203
LOCUS AF317203 10972 bp RNA linear VRL 11-FEB-2001
DEFINITION West Nile virus VLG-4 polyprotein precursor, gene, complete cds.
ACCESSION AF317203
VERSION AF317203.1 GI:12744408
KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE 1 (bases 1 to 10972)

AUTHORS Platonov,A.E., Karan,L., Yazishina,S., Obukhov,I.L., Shipulina,O.
and Shipulin,G.A.
TITLE Genetic similarity of West Nile viruses caused epidemics in
Volgograd 1999 and Romania 1996
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 10972)
AUTHORS Karan,L., Yazishina,S., Obukhov,I.L., Shipulina,O., Shipulin,G.A.
and Platonov,A.E.
TITLE Direct Submission
JOURNAL Submitted (26-OCT-2000) Central Research Institute of Epidemiology,
Novogireevskaya Str. 3A, Moscow 111123, Russia
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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 10972;
Best Local Similarity 100.0%; Pred. No. 1.6; Mismatches 0; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CCGGCTGTCAATATGCTAAA 21
|||||
Db 97 CCGGCTGTCAATATGCTAAA 117
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RESULT 9
AF206518
LOCUS AF206518 10975 bp DNA linear VRL 08-MAY-2000
DEFINITION West Nile virus isolate 2741, complete genome.
ACCESSION AF206518
VERSION AF206518.2 GI:7717200
KEYWORDS
SOURCE West Nile virus
ORGANISM West Nile virus

Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
REFERENCE 1 (bases 1 to 10975)
Anderson,J.F., Andreadis,T.G., Vossbrinck,C.R., Tirrell,S.,
Wakem,E.M., French,R.A., Garmendia,A.E. and Van Kruiningen,H.J.
Isolation of West Nile virus from mosquitoes, crows, and a Cooper's
hawk in Connecticut
JOURNAL Science 286 (5448), 2331-2333 (1999)
MEDLINE 20070287
PUBMED 10600741

REFERENCE 2 (bases 1 to 10975)
Vossbrinck,C.R., Anderson,J.F. and Andreadis,T.G.
AUTHORS Genome Sequence of West Nile Virus from Culex pipiens isolate
TITLE Unpublished
JOURNAL
REFERENCE 3 (bases 1 to 10975)
Anderson,J.F., Andreadis,T.G. and Vossbrinck,C.R.
AUTHORS Direct Submission
TITLE Submitted (18-NOV-1999) Soil and Water, Connecticut Agricultural

Experiment Station, 123 Huntington Street, New Haven, CT 06511, USA
4 (bases 1 to 10975)
Anderson,J.F., Andreadis,T.G. and Vossbrinck,C.R.
Direct Submission
Submitted (08-May-2000) Soil and Water, Connecticut Agricultural
Experiment Station, 123 Huntington Street, New Haven, CT 06511, USA
Sequence update by submitter
COMMENT On May 8, 2000 this sequence version replaced gi:6636507.
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ORIGIN

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Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 111 CCGGGCTGTCAATATGCTAAA 131

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AY262283 LOCUS AY262283 10984 bp RNA linear VRL 29-OCT-2003

DEFINITION West Nile virus isolate KN3829 polyprotein gene, complete cds.

ACCESSION AY262283

VERSION AY262283.1 GI:30230630

KEYWORDS

SOURCE West Nile virus (WNV)

ORGANISM Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

Flavivirus; Japanese encephalitis virus group.

REFERENCE 1 (bases 1 to 10984)

AUTHORS Charrat, R., Brault, A.C., Gallian, P., Lemasson, J.-J., Murgue, B.,

Murri, S., Paetorino, B., Zeller, H., de Chesse, R., de Micco, P. and de

Lamballerie, X.

TITLE Evolutionary relationship between Old World West Nile virus

strains. Evidence for viral gene flow between africa, the middle

east, and europe

JOURNAL Virology 315 (2), 381-388 (2003)

MEDLINE 22949215

PUBMED 14585341

REFERENCE 2 (bases 1 to 10984)

AUTHORS Brault, A.C. and de Lamballerie, X.

TITLE Direct Submission

JOURNAL Submitted (25-MAR-2003) Division of Vector-Borne Infectious

Diseases, Centers for Disease Control and Prevention, P.O. Box

2087, Fort Collins, CO 80522, USA

FEATURES Location/Qualifiers

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ORIGIN

Query Match

100.0%; Score 21; DB 14; Length 10984;

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Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGCTCAATATGCTAAA 21
Db 93 CCGGGCTGCTCAATATGCTAAA 113

RESULT 11
AY268132 West Nile virus strain PaAn001 polyprotein (pol) gene, complete
LOCUS AY268132 10989 bp RNA linear VRL 03-NOV-2003
DEFINITION AY268132.1 GI:33242574
VERSION AY268132.1
KEYWORDS West Nile virus (WNV)
SOURCE West Nile virus
ORGANISM West Nile virus
VIRUSES; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 10989)
REFERENCE Charrel,R.N., Brault,A.C., Gallian,P., Lemasson,J.-J., Murgue,B.,
AUTHORS Murri,S., Pastorino,B., Zeller,H., de chesse,R., de Micco,P. and de
Lamballerie,X.
TITLE Evolutionary relationship between Old World West Nile virus
strains. Evidence for viral gene flow between africa, the middle
east, and europe
JOURNAL Virology 315 (2), 381-388 (2003)
MEDLINE 22949215
PUBMED 14585341
REFERENCE de Lamballerie,X., Brault,A.C., Gallian,P., Lemasson,J., Murgue,B.,
AUTHORS Murri,S., Pastorino,B., Zeller,H., Dechesse,R., de Micco,P. and
Charrel,R.N.
TITLE Direct Submision
JOURNAL Submitted (03-APR-2003) Virology, Medical University, 27 bd Jean
JOURNAL Moulin, Marseille 13005, France
FEATURES
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Query Match 100.0%; Score 21; DB 14; Length 10989;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CCGGGCTGCTCAATATGCTAAA 21
Db 109 CCGGGCTGCTCAATATGCTAAA 129

RESULT 12
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LOCUS AY268133 10989 bp RNA linear VRL 03-NOV-2003
DEFINITION West Nile virus strain PaH001 polyprotein (pol) gene, complete cds.
ACCESSION AY268133
VERSION AY268133.1 GI:33242576
KEYWORDS West Nile virus (WNV)
SOURCE West Nile virus
ORGANISM West Nile virus
VIRUSES; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 (bases 1 to 10989)
REFERENCE Charrel,R.N., Brault,A.C., Gallian,P., Lemasson,J.-J., Murgue,B.,
AUTHORS Murri,S., Pastorino,B., Brault,A.C., Zeller,H., de chesse,R., de Micco,P. and de
Lamballerie,X.
TITLE Evolutionary relationship between Old World West Nile virus
strains. Evidence for viral gene flow between africa, the middle
east, and europe
JOURNAL Virology 315 (2), 381-388 (2003)
MEDLINE 22949215
PUBMED 14585341
REFERENCE de Lamballerie,X., Brault,A.C., Gallian,P., Lemasson,J., Murgue,B.,
AUTHORS Murri,S., Pastorino,B., Zeller,H., Dechesse,R., de Micco,P. and
Murri,S., Pastorino,B., Zeller,H., de chesse,R., de Micco,P. and

[illegible]

Charrel, R.N.
Direct Submission
Submitted (03-APR-2003) Virology, Medical University, 27 bd Jean
Moulin, Marseille 13005, France

FEATURES
source

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ORIGIN

Query Match 100.0%; Score 21; DB 14; Length 10989;
Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0

Qy 1 CCGGGCTGTCATATGCTAAA 21
|||||

Db 109 CCGGGCTGTCATATGCTAAA 129
|||||

RESULT 13
AY274504

LOCUS
AY274504 11022 bp mRNA linear VRL 02-JUL-2003
DEFINITION
Kunjin virus clone FLSDX polyprotein mRNA, complete cds.
ACCESSION
AY274504
VERSION
AY274504.1 GI:32306849

KEYWORDS
Kunjin virus

SOURCE
Kunjin virus

ORGANISM
Kunjin virus

REFERENCE
1 (bases 1 to 11022)
Viruses; sRNA positive-strand viruses, no DNA stage; Flaviviridae
Flavivirus; Japanese encephalitis virus group.

AUTHORS
1 (bases 1 to 11022)
Liu W.J., Chen, H.B. and Khromykh, A.A.

TITLE
Molecular and Functional Analyses of Kunjin Virus Infectious cDNA
Clones Demonstrate the Essential Roles for NS2A in Virus Assembly
and for a Nonconservative Residue in NS3 in RNA Replication

JOURNAL
J. Virol. 77 (14), 7804-7813 (2003)

MEDLINE
22713678

PUBMED
12829820

REFERENCE
2 (bases 1 to 11022)
Khromykh, A.A., Liu, W.J. and Chen, H.B.
Direct Submission
Submitted (11-APR-2003) Clinical Medical Virology Centre,
University of Queensland/Sir Albert Sakzewski Virus Research
Centre, Royal Children's Hospital, Herston Rd., Herston, Brisbane,
QLD 4029, Australia

FEATURES
source

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Best Local Similarity 100.0%; Pred. No. 1.6; Mismatches 0; Gaps 0;
Matches 21; Conservative 0; Indels 0; Gaps 0;
Qy 1 CCGGGCTGTCAATATGCTAAA 21
Db 129 CCGGGCTGTCAATATGCTAAA 149
RESULT 14
AY274505
LOCUS AY274505 11022 bp mRNA linear VRL 02-JUL-2003
DEFINITION Kunjin virus clone pAKUN polyprotein mRNA, complete cds.
ACCESSION AY274505
VERSION AY274505.1 GI:32306851
KEYWORDS
SOURCE Kunjin virus
ORGANISM Kunjin virus
REFERENCE 1 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
2 (bases 1 to 11022)
AUTHORS Liu, W.J., Chen, H.B. and Khromykh, A.A.
TITLE Molecular and Functional Analyses of Kunjin Virus Infectious cDNA
Clones Demonstrate the Essential Roles for NS2A in Virus Assembly
and for a Nonconservative Residue in NS3 in RNA Replication
J. Virol. 77 (14), 7804-7813 (2003)
JOURNAL 22713678
MEDLINE 12829820
PUBMED 2 (bases 1 to 11022)
REFERENCE Khromykh, A.A., Liu, W.J. and Chen, H.B.
AUTHORS Direct Submission
TITLE Submitted (11-APR-2003) Clinical Medical Virology Centre,
University of Queensland/Sir Albert Sakzewski Virus Research
Centre, Royal Children's Hospital, Herston Rd., Herston, Brisbane,
QLD 4029, Australia
FEATURES
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VERSION AY490240.2 GI:46277828
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ORGANISM West Nile virus
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REFERENCE 1 (bases 1 to 11028)
AUTHORS Jiang, T., Qin, E. and Deng, Y.
TITLE Sequence determination and analysis of West Nile Virus strain
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 11028)
AUTHORS Jiang, T., Qin, E. and Deng, Y.
TITLE Direct Submission
JOURNAL Submitted (28-NOV-2003) Virology, Institute of Microbiology and
Epidemiology, Fengtai Dongda Street, Beijing 100071, China
REFERENCE 3 (bases 1 to 11028)
AUTHORS Jiang, T., Qin, E. and Deng, Y.
TITLE Direct Submission
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REMARK
COMMENT On Apr 8, 2004 this sequence version replaced gi:40362614.
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DEFINITION Sequence 1 from Patent WO02081741.
ACCESSION AX577796
VERSION AX577796.1 GI:27647035
KEYWORDS
SOURCE
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Flavivirus sp.
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus.

REFERENCE
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Guenet, J.L., Moshimo, T., Simon-Chazottes, D., Montagnetelli, X.,
Frankel, M.P., Despres, P., Deubel, V., Bonhomme, F., and Lucas, M.
Use of products of genes of the 2', 5' oligoadenylate synthetase
family (oas) for screening antiviral agents and for detecting
responsiveness to flaviviridae infection
Patent: WO 02081741-A 1 17-OCT-2002;
INSTITUT PASTEUR (FR); CENTRE NATIONAL DE LA RECHERCHE
SCIENTIFIQUE (CNRS) (FR)

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ABI85914
VERSION
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KEYWORDS
SOURCE
West Nile virus (WNV)
ORGANISM
West Nile virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.

REFERENCE
1
Shirato, K., Miyoshi, H., Goto, A., Ako, Y., Ueki, T., Kariwa, H. and
Takahama, I.
Correlation between viral envelope glycosylation and
neuroinvasiveness of the New York strain of the West Nile virus
Unpublished
Shirato, K., Kariwa, H. and Takahama, I.
Direct Submission
Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of
Veterinary Medicine, Hokkaido University, Laboratory of Public
Health, Department of Environmental Veterinary Medicine; Kita-19
Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
(E-mail: shirato@vetmed.hokudai.ac.jp, Tel: 81-11-706-5213),
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On Jul 30, 2004 this sequence version replaced gi:50838778.

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ORIGIN

| | | | | |
|-----------------------|-----------------|----------------|-----------|---------------|
| Query Match | 100.0%; | Score 21; | DB 14; | Length 11029; |
| Best Local Similarity | 100.0%; | Pred. No. 1.6; | | |
| Matches 21; | Conservative 0; | Mismatches 0; | Indels 0; | Gaps 0; |

| | | | |
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| Qy | 1 | CCGGGCTGCTCAATATGCTAAA | 21 |
| Db | 129 | CCGGGCTGCTCAATATGCTAAA | 149 |

| | |
|-------------|--|
| RESULT 19 | |
| AB185915 | |
| LOCUS | |
| DEFINITION | AB185915 11029 bp RNA linear VRL 31-JUL-2004 |
| | West Nile virus gene for polyprotein precursor protein, complete |
| | cds, isolate: 6-Sp. |
| ACCESSION | AB185915 |
| VERSION | AB185915.2 |
| GI:50872125 | |
| KEYWORDS | |
| SOURCE | West Nile virus (WNV) |
| ORGANISM | West Nile virus |

Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Flavivirus; Japanese encephalitis virus group.
1 Shirato, K., Miyoshi, H., Goto, A., Ako, Y., Ueki, T., Kariwa, H. and Takashima, I.
Correlation between viral envelope glycosylation and neuroinvasiveness of the New York strain of the West Nile virus
Unpublished
2 (bases 1 to 11029)
Shirato, K., Kariwa, H. and Takashima, I.
Direct Submission
Submitted (28-JUL-2004) Kazuya Shirato, Graduate School of Veterinary Medicine, Hokkaido University, Laboratory of Public Health, Department of Environmental Veterinary Medicine; Kita-19 Nishi-9, Kita-ku, Sapporo, Hokkaido 060-0818, Japan
(E-mail: shirato@vetmed.hokudai.ac.jp, Tel: 81-11-706-5213 (ex. 5213), Fax: 81-11-706-5213)
On Jul 30, 2004 this sequence version replaced gi:50838780.

COMMENT FEATURES

source

Location/Qualifiers

1. 11029

/organism="West Nile virus"

/mol_type="genomic RNA"

/strain="NY99-6922"

/isolate="6-Sp"

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ORIGIN

```
Query Match      100.0%; Score 21; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. NO. 1.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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| | | | | | |
|------------|---|-------------|-----|--------|-----------------|
| RESULT 20 | AB185916 | 11029 bp | RNA | linear | VRL 30-JUL-2004 |
| LOCUS | AB185916 | | | | |
| DEFINITION | West Nile virus gene for polyprotein precursor protein, complete cds, isolate: B-SP. | | | | |
| ACCESSION | AB185916 | | | | |
| VERSION | AB185916.1 | GI:50838782 | | | |
| KEYWORDS | | | | | |
| SOURCE | West Nile virus (WNV) | | | | |
| ORGANISM | West Nile virus Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Flavivirus; Japanese encephalitis virus group. | | | | |

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ORIGIN

| | | | | |
|-----------------------|-----------------|----------------|-----------|---------------|
| Query Match | 100.0% | Score 21; | DB 14; | Length 11029; |
| Best Local Similarity | 100.0% | Pred. NO. 1.6; | | |
| Matches 21; | Conservative 0; | Mismatches 0; | Indels 0; | Gaps 0; |

RESULT 21
AB185917
LOCUS AB185917 11029 bp RNA linear VRL 30-JUL-2004
DEFINITION West Nile virus gene for polyprotein precursor protein, complete cds, isolate: B-LP.
ACCESSION AB185917
VERSION AB185917.1 GI:50838784
KEYWORDS


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6916..7680      /product="non-structural protein NS4B"
7681..10395      /product="non-structural protein NS5"
ORIGIN
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Query Match      100.0%; Score 21; DB 14; Length 11029;
Best Local Similarity 100.0%; Pred. No. 1.6; Mismatches 0; Indels 0; Gaps 0;
Matches 21; Conservative 0;

Qy      1 CCGGGCTGCTCAATATGCTAAA 21
      |||||
Db      129 CCGGGCTGCTCAATATGCTAAA 149
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Search completed: September 6, 2005, 20:29:40
Job time : 740.656 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 16:01:23 ; Search time 189.656 Seconds
(without alignments)
655.473 Million cell updates/sec

Title: US-10-729-421-34

Perfect score: 21

Sequence: 1 ccgggtgtcaatgctaaa 21

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database :

N_Geneseq_16Dec04:*

- 1: Geneseqn1980s:*
- 2: Geneseqn1990s:*
- 3: Geneseqn2000s:*
- 4: Geneseqn2001as:*
- 5: Geneseqn2001bs:*
- 6: Geneseqn2002as:*
- 7: Geneseqn2002bs:*
- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
- 10: Geneseqn2003cs:*
- 11: Geneseqn2003ds:*
- 12: Geneseqn2004as:*
- 13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|--------------------|
| 1 | 21 | 100.0 | 21 | 12 | ADQ30664 West Nile |
| 2 | 21 | 100.0 | 37 | 12 | ADN36773 West Nile |
| 3 | 21 | 100.0 | 65 | 12 | ADN36771 West Nile |
| 4 | 21 | 100.0 | 365 | 6 | ABK51710 Partial c |
| 5 | 21 | 100.0 | 366 | 8 | ABQ76684 WNVcwt DN |
| 6 | 21 | 100.0 | 967 | 12 | ADQ30647 West Nile |
| 7 | 21 | 100.0 | 10945 | 13 | ADR32078 Genomic D |
| 8 | 21 | 100.0 | 10945 | 13 | ADR67768 West Nile |
| 9 | 21 | 100.0 | 10962 | 12 | ADK13681 West Nile |
| 10 | 21 | 100.0 | 10975 | 12 | ADN98022 West Nile |
| 11 | 21 | 100.0 | 11029 | 8 | ABZ68481 Nucleotid |
| 12 | 21 | 100.0 | 11029 | 10 | ABV74821 West Nile |
| 13 | 21 | 100.0 | 11029 | 12 | ADN98023 West Nile |
| 14 | 19 | 90.5 | 21 | 12 | ADN36693 West Nile |
| 15 | 19 | 90.5 | 24 | 12 | ADN36823 West Nile |
| 16 | 19 | 90.5 | 48 | 12 | ADN36707 West Nile |
| 17 | 19 | 90.5 | 69 | 12 | ADN36694 West Nile |
| 18 | 18 | 85.7 | 20 | 12 | ADN36696 West Nile |
| 19 | 18 | 85.7 | 47 | 12 | ADN36708 West Nile |
| 20 | 17.8 | 84.8 | 4512 | 2 | AAQ22767 JEV Nakay |

| | | | | | | |
|----|------|------|--------|----|----------|---------------------|
| 21 | 17.8 | 84.8 | 10818 | 12 | ADO07431 | Ado07431 Japanese |
| 22 | 17.8 | 84.8 | 10968 | 12 | ADO07437 | Ado07437 Japanese |
| 23 | 17.8 | 84.8 | 10976 | 3 | ABL50890 | Abi50890 Japanese |
| 24 | 17.8 | 84.8 | 18563 | 12 | ADO07466 | Ado07466 Japanese |
| 25 | 17.8 | 84.8 | 18563 | 12 | ADO07465 | Ado07465 Japanese |
| 26 | 17.8 | 84.8 | 18565 | 12 | ADO07467 | Ado07467 Japanese |
| 27 | 17.8 | 84.8 | 19038 | 12 | ADO07468 | Ado07468 Japanese |
| 28 | 17.8 | 84.8 | 19038 | 12 | ADO07469 | Ado07469 Japanese |
| 29 | 17.8 | 84.8 | 19040 | 12 | ADO07470 | Ado07470 Japanese |
| 30 | 17 | 81.0 | 17 | 6 | ACN01436 | Acn01436 WNV Inozy |
| 31 | 17 | 81.0 | 17 | 6 | ACN00029 | Acn00029 WNV Hamme |
| 32 | 17 | 81.0 | 17 | 6 | ACN09480 | Acn09480 WNV minus |
| 33 | 17 | 81.0 | 17 | 6 | ACN15271 | Acn15271 WNV minus |
| 34 | 17 | 81.0 | 17 | 6 | ACN14165 | Acn14165 WNV minus |
| 35 | 17 | 81.0 | 17 | 6 | ACN04724 | Acn04724 WNV DNazY |
| 36 | 17 | 81.0 | 17 | 6 | ACN09479 | Acn09479 WNV minus |
| 37 | 17 | 81.0 | 20 | 12 | ADN36776 | Adn36776 West Nile |
| 38 | 16.8 | 80.0 | 2262 | 8 | ACA54320 | Acas4320 Prokaryot |
| 39 | 16.4 | 78.1 | 356 | 9 | ACH31834 | Ach31834 Human bon |
| 40 | 16.2 | 77.1 | 326 | 6 | ABN75146 | Abn75146 Human ORF |
| 41 | 16.2 | 77.1 | 1149 | 10 | ADB46059 | Adb46059 rscP DNA |
| 42 | 16.2 | 77.1 | 2023 | 10 | ADB69062 | Adb69062 C. neofor |
| 43 | 16.2 | 77.1 | 2232 | 11 | ACH98107 | Ach98107 Klebsiell |
| 44 | 16.2 | 77.1 | 3177 | 4 | ABL16411 | Abi16411 Drosophil |
| 45 | 16.2 | 77.1 | 7018 | 4 | ABL16410 | Abi16410 Drosophil |
| 46 | 16 | 76.2 | 17 | 6 | ACN00030 | Acn00030 WNV Hamme |
| 47 | 16 | 76.2 | 17 | 6 | ACN03477 | Acn03477 WNV ZinzY |
| 48 | 16 | 76.2 | 2005 | 4 | AAF81805 | Aaf81805 Human sec |
| 49 | 16 | 76.2 | 149671 | 6 | ABK84797 | Abk84797 Human cDN |
| 50 | 16 | 76.2 | 149671 | 9 | ADB70361 | Adb70361 Moezin cD |
| 51 | 16 | 76.2 | 149671 | 12 | ADJ37140 | Adj37140 Human mal |
| 52 | 15.8 | 75.2 | 624 | 6 | ABN73108 | Abn73108 Bovine em |
| 53 | 15.8 | 75.2 | 1197 | 5 | AAS81962 | Aas81962 DNA encod |
| 54 | 15.8 | 75.2 | 1208 | 5 | AAS74882 | Aas74882 DNA encod |
| 55 | 15.8 | 75.2 | 1208 | 5 | AAS93301 | Aas93301 DNA encod |
| 56 | 15.8 | 75.2 | 1208 | 5 | AAS77346 | Aas77346 DNA encod |
| 57 | 15.8 | 75.2 | 2373 | 5 | AAS86894 | Aas86894 DNA encod |
| 58 | 15.8 | 75.2 | 11184 | 12 | ADP86274 | Adp86274 Hepatitis |
| 59 | 15.8 | 75.2 | 11184 | 12 | ADP86276 | Adp86276 Hepatitis |
| 60 | 15.8 | 75.2 | 11184 | 12 | ADP86277 | Adp86277 Hepatitis |
| 61 | 15.8 | 75.2 | 11313 | 12 | ADP86273 | Adp86273 Hepatitis |
| 62 | 15.8 | 75.2 | 11313 | 12 | ADP86264 | Adp86264 Hepatitis |
| 63 | 15.8 | 75.2 | 11313 | 12 | ADP86266 | Adp86266 Hepatitis |
| 64 | 15.8 | 75.2 | 11313 | 12 | ADP86265 | Adp86265 Hepatitis |
| 65 | 15.8 | 75.2 | 11313 | 12 | ADP86268 | Adp86268 Hepatitis |
| 66 | 15.8 | 75.2 | 11313 | 12 | ADP86270 | Adp86270 Hepatitis |
| 67 | 15.8 | 75.2 | 11313 | 12 | ADP86271 | Adp86271 Hepatitis |
| 68 | 15.8 | 75.2 | 11313 | 12 | ADP86272 | Adp86272 Hepatitis |
| 69 | 15.8 | 75.2 | 11313 | 12 | ADP86269 | Adp86269 Hepatitis |
| 70 | 15.8 | 75.2 | 11313 | 12 | ADP86275 | Adp86275 Hepatitis |
| 71 | 15.8 | 75.2 | 11313 | 12 | ADP86267 | Adp86267 Hepatitis |
| 72 | 15.8 | 75.2 | 12306 | 10 | ADI41414 | Adi41414 BB7 nucle |
| 73 | 15.8 | 75.2 | 12315 | 10 | ADI41413 | Adi41413 BB7M4bRLu |
| 74 | 15.8 | 75.2 | 12980 | 2 | AAV59364 | Aav59364 Hepatitis |
| 75 | 15.8 | 75.2 | 12980 | 6 | ABK87286 | Abk87286 Hepatitis |
| 76 | 15.8 | 75.2 | 12980 | 8 | ACA62469 | Acac62469 DNA encod |
| 77 | 15.8 | 75.2 | 15065 | 3 | AAZ36195 | Aaz36195 Nucleotid |
| 78 | 15.8 | 75.2 | 16847 | 12 | ADO07464 | Ado07464 Japanese |
| 79 | 15.4 | 73.3 | 444 | 9 | ACH24983 | Ach24983 Human adu |
| 80 | 15.4 | 73.3 | 1119 | 6 | AAD31757 | Adad31757 Soybean H |
| 81 | 15.4 | 73.3 | 73583 | 12 | ADQ59187 | Adq59187 MSI-H car |
| 82 | 15.4 | 73.3 | 22930 | 6 | ABK84349 | Abk84349 Human cDN |
| 83 | 15.4 | 73.3 | 295096 | 11 | ACN44068 | Acn44068 Mouse gen |
| 84 | 15.2 | 72.4 | 26 | 12 | ADN36839 | Adn36839 West Nile |
| 85 | 15.2 | 72.4 | 445 | 4 | AAS36425 | Aas36425 Human car |
| 86 | 15.2 | 72.4 | 445 | 10 | ADS47119 | Ads47119 Human car |
| 87 | 15.2 | 72.4 | 445 | 13 | ADJ08537 | Adj08537 Human car |
| 88 | 15.2 | 72.4 | 452 | 3 | AAA82296 | Aaa82296 N. mening |
| 89 | 15.2 | 72.4 | 914 | 5 | AAS56345 | Aas56345 DNA encod |
| 90 | 15.2 | 72.4 | 1038 | 3 | AZ60397 | Az60397 A diacylg |
| 91 | 15.2 | 72.4 | 1065 | 12 | ADP72956 | Adp72956 Renal tox |
| 92 | 15.2 | 72.4 | 1121 | 2 | AAQ21554 | Aaq21554 Polyfunct |
| 93 | 15.2 | 72.4 | 1121 | 6 | ABK63744 | Abk63744 Rat seque |

94 15.2 72.4 1121 10 ADB58300 Adb58300 Toxicity-
 95 15.2 72.4 1121 10 ADB52851 Adb52851 Primary r
 96 15.2 72.4 1121 10 ABT41990 Abt41990 Toxicity
 97 15.2 72.4 1167 10 ADF01035 Adf01035 Bacterial
 c 98 15.2 72.4 1302 4 AAF58403 Aaf58403 Rat rOCIL
 c 99 15.2 72.4 1389 11 ACH96542 Ach96542 Klebsiell
 100 15.2 72.4 1418 6 ABZ15428 Abz15428 Arabidops

ALIGNMENTS

RESULT 1

ADQ30664

ID ADQ30664 standard; DNA; 21 BP.

XX

AC ADQ30664;

XX

DT 23-SEP-2004 (first entry)

XX

DE West Nile Virus capsid gene sense primer WNVV1.

XX

KW ss; primer; West Nile Virus; diagnosis.

XX

OS West Nile virus.

XX

FN WO2004055159-A2.

XX

XX

PD 01-JUL-2004.

XX

PF 05-DEC-2003; 2003WO-US038750.

XX

PR 12-DEC-2002; 2002US-0432850P.

XX

PR 20-JUN-2003; 2003US-0480431P.

XX

XX (CHIR) CHIRON CORP.

PA

PI Shyamala V;

XX

XX WPI; 2004-488058/46.

DR

PT New isolated oligonucleotides for accurately diagnosing West Nile virus

XX

PT infection or for capturing, detecting and quantitating West Nile virus in

XX

PT blood samples.

XX

PS Claim 1; SEQ ID NO 34; 56pp; English.

XX

CC The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to a PCR primer to amplify a fragment of the capsid gene of
 CC the WNV genome. The fragment is detected using the oligonucleotides of
 CC the invention.

XX Sequence 21 BP; 6 A; 5 C; 5 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 21;

Best Local Similarity 100.0%; Pred. No. 0.38;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21

|||||

Db 1 CCGGGCTGTCATATGCTAAA 21

RESULT 2

ADN36773

ID ADN36773 standard; DNA; 37 BP.

XX

AC ADN36773;

XX

DT 15-JUL-2004 (first entry)

XX

DE West Nile virus detection-related oligonucleotide probe SeqID95.

XX

KW hybridisation assay probe; nucleic acid detection;

XX

KW target-complementary sequence; flavivirus; West Nile virus; WNV;

XX

KW RNA virus; infection; meningitis; encephalitis;

XX

KW high throughput screening; probe; ss.

XX

OS West Nile virus.

XX

FN WO2004036190-A2.

XX

XX 29-APR-2004.

XX

XX 10-OCT-2003; 2003WO-US033639.

XX

XX 16-OCT-2002; 2002US-0418891P.

XX

XX 25-NOV-2002; 2002US-0429006P.

XX

XX 24-FEB-2003; 2003US-0449810P.

XX

XX (GENP-) GEN-PROBE INC.

PA

PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX

XX WPI; 2004-389590/36.

DR

PT New hybridization assay probe comprising target-complementary sequence of
 PT bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Claim 55; SEQ ID NO 95; 135pp; English.

XX

CC This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.

XX Sequence 37 BP; 10 A; 10 C; 11 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 37;

Best Local Similarity 100.0%; Pred. No. 0.41;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21

|||||

Db 4 CCGGGCTGTCATATGCTAAA 24

RESULT 3

ADN36771

ID ADN36771 standard; DNA; 65 BP.

XX

AC ADN36771;

XX

XX 15-JUL-2004 (first entry)

DT

DE West Nile virus detection-related oligonucleotide probe SeqID93.
 XX hybridisation assay probe; nucleic acid detection;
 KW target-complementary sequence; flavivirus; West Nile virus; WNV;
 KW RNA virus; infection; meningitis; encephalitis;
 KW high throughput screening; probe; ss.

OS West Nile virus.

XX WO2004036190-A2.

PN 29-APR-2004.

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

PR 25-NOV-2002; 2002US-0429006P.

PR 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JW, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of

PT bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Example 6; SEQ ID NO 93; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
 CC detecting a nucleic acid, which is a probe sequence that comprises a
 CC target-complementary sequence of bases, and optionally one or more base
 CC sequences that are not complementary to the nucleic acid that is to be
 CC detected. The hybridisation assay probes and the kits are useful in
 CC detecting and amplifying a target nucleic acid sequence, for example
 CC flavivirus like West Nile virus, that may be present in a biological
 CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
 CC birds and culex mosquitoes, with humans and horses serving as incidental
 CC hosts. Infection of humans can lead to meningitis or encephalitis. The
 CC invention may allow for accurate and efficient high throughput screening.
 CC The present sequence is that of an oligonucleotide probe which is related
 CC to the invention.

XX Sequence 65 BP; 19 A; 17 C; 20 G; 9 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 65;

Best Local Similarity 100.0%; Pred. No. 0.44; Mismatches 0; Indels 0; Gaps 0;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCAATATGCTAAA 21

Db 32 CCGGGCTGTCAATATGCTAAA 52

RESULT 4

ABK51710

ID ABK51710 standard; cDNA; 365 BP.

XX ABK51710;

XX 27-AUG-2002 (first entry)

XX Partial cDNA for west nile virus capsid protein.

XX Human; ss, IgE leader sequence; west nile virus capsid protein;
 KW RNA secondary structure; free energy; gene therapy; cancer;
 KW hyperproliferative disease; autoimmune disease; rheumatoid arthritis;
 KW multiple sclerosis; Sjogren's syndrome; sarcoidosis; scleroderma;
 KW insulin-dependent diabetes mellitus; autoimmune thyroiditis; psoriasis;
 KW reactive arthritis; ankylosing spondylitis; polymyositis; vasculitis;
 KW dermatomyositis; Crohn's disease; ulcerative colitis.

XX

OS West Nile virus.

XX WO200229088-A2.

XX 11-APR-2002.

XX 04-OCT-2001; 2001WO-US031451.

XX 04-OCT-2000; 2000US-0237885P.

XX (UYPE-) UNIV PENNSYLVANIA.

XX Weiner DB, Yang J;

XX WPI; 2002-416682/44.

XX Producing recombinant protein for preparing pharmaceutical compounds to
 CC treat, e.g., cancers or autoimmune disorders, comprises predicting
 CC secondary structure (SS) of mRNA and modifying DNA to give mRNA with SS
 CC having increased free energy.

XX Example 2; Fig 1; 48pp; English.

XX The invention relates to producing (M1) a protein (I) in a recombinant
 CC expression system (II) comprising: (a) predicting the secondary structure
 CC of mRNA; (b) modifying the native heterologous DNA sequence where the
 CC mRNA transcribed from the modified DNA has a secondary structure with
 CC increased free energy; and (c) using the modified DNA in (II) for
 CC production of (I). Also included are (1) an injectable pharmaceutical
 CC composition comprising a nucleic acid molecule that includes a modified
 CC coding sequence (IV) encoding a protein operably linked to regulatory
 CC elements, where (IV) comprises a higher AT or AU content relative to the
 CC AT or AU content of the native coding sequence and further comprising a
 CC pharmaceutical carrier and (2) a recombinant viral vector comprising a
 CC nucleic acid molecule that includes (IV). The method is used for
 CC producing a protein in a recombinant expression system. Use of a nucleic
 CC acid or recombinant viral vector that have modified DNA sequences to
 CC improve protein production can be used in gene therapy and for the
 CC treatment of cancers, hyperproliferative diseases, and autoimmune
 CC diseases such as rheumatoid arthritis, multiple sclerosis, Sjogren's
 CC syndrome, sarcoidosis, insulin-dependent diabetes mellitus, scleroderma,
 CC thyroiditis, reactive arthritis, ankylosing spondylitis, Crohn's disease and
 CC polymyositis, dermatomyositis, psoriasis, vasculitis, ulcerative colitis.
 CC The present sequence is a cDNA for West Nile virus
 CC capsid protein. Fusion constructs of modified mRNA for the capsid protein
 CC and human IgE leader sequence are used in an experiment to minimise the
 CC free energy of the capsid protein mRNA. Note: The present sequence is not
 CC shown in the specification but was created using the information in
 CC figure 1 and the sequence appearing as ABK51708

XX Sequence 365 BP; 103 A; 80 C; 109 G; 73 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 6; Length 365;

Best Local Similarity 100.0%; Pred. No. 0.57; Mismatches 0; Indels 0; Gaps 0;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCAATATGCTAAA 21

Db 29 CCGGGCTGTCAATATGCTAAA 49

RESULT 5

ABQ76684

ID ABQ76684 standard; DNA; 366 BP.

XX ABQ76684;

XX 13-MAY-2003 (first entry)

XX WNVcwt DNA fragment.

XX Capsid protein; WNVcwt; mRNA secondary structure; cancer; immunosuppressive; antirheumatic; cytostatic; antitumor; neuroprotective;

KW antiarthritic; antidiabetic; antithyroid; antipsoriatic; virucide; gene;
 KW antiparasitic; antiallergic; gene therapy; allergen; multiple sclerosis;
 KW protective immune response; hyperproliferative cell; ulcerative colitis;
 KW hyperproliferative disease; psoriasis; autoimmune disease; psoriasis;
 KW rheumatoid arthritis; Sjogren's syndrome; autoimmune thyroiditis;
 KW insulin dependent diabetes mellitus; Crohn's disease; ds.

XX West Nile virus.

XX Key Location/Qualifiers
 FT CDS 1..366
 FT /*tag= a
 FT /product= "WNVCwt"
 FT /note= "no start or stop codon given"

XX US2002123099-A1.

XX 05-SEP-2002.

XX 04-OCT-2001; 2001US-00971806.

XX 04-OCT-2000; 2000US-0237885P.

XX (WEIN/) WEINER D B.
 XX (YANG/) YANG J.

XX Weiner DB, Yang J;

XX WPI; 2003-066795/06.
 XX P-PSDB; ABG73556.

XX Producing protein in recombinant expression system involves predicting
 PT secondary structure of RNA encoding a protein and increasing free energy
 PT for the secondary structure by modifying sequence of DNA encoding the
 PT RNA.

XX Example 2; Fig 1; 25pp; English.

XX This invention describes a novel method for producing a protein by
 CC translation of mRNA from heterologous DNA sequences. The method involves
 CC predicting the secondary structure of mRNA transcribed from a native
 CC heterologous DNA sequence, modifying the sequence where mRNA transcribed
 CC from the modified DNA sequence has a secondary structure with increased
 CC free energy compared to mRNA transcribed from native DNA and using
 CC modified heterologous DNA for protein production. The products of the
 CC invention have immunosuppressive, antirheumatic, cytostatic, antiulcer,
 CC neuroprotective, antiarthritic, antidiabetic, antithyroid, antipsoriatic,
 CC virucide, antiparasitic and anti-allergic activity and can be used for
 CC gene therapy. The method described is useful for producing a protein in a
 CC recombinant expression system, preferably a cell free in vitro
 CC transcription and translation system, an in vitro cell expression system,
 CC a DNA construct used in direct DNA injection, or a recombinant vector for
 CC delivery of DNA to an individual. The products of the invention are
 CC useful for eliciting broad immune responses against a target protein,
 CC i.e. proteins specifically associated with pathogens such as viruses,
 CC parasites, allergens, or the individual's own abnormal cells.
 CC Compositions containing the products of the invention confer a broad
 CC based protective immune response against hyperproliferative cells that
 CC are characteristic in hyperproliferative diseases including all forms of
 CC cancer and psoriasis. Such compositions are also useful for treating
 CC individuals suffering from autoimmune diseases including rheumatoid
 CC arthritis, multiple sclerosis, Sjogren's syndrome, insulin dependent
 CC diabetes mellitus, autoimmune thyroiditis, Crohn's disease, ulcerative
 CC colitis and psoriasis. This sequence encodes the West Nile virus wild-
 CC type capsid protein described as WNVCwt in the disclosure of the
 CC invention

XX Sequence 366 BP; 103 A; 81 C; 108 G; 74 T; 0 U; 0 Other;

XX Query Match 100.0%; Score 21; DB 8; Length 366;
 XX Best Local Similarity 100.0%; Pred. No. 0.57; Length 366;
 XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
 |||||
 Db 30 CCGGGCTGTCATATGCTAAA 50

RESULT 6

ID ADQ30647 standard; DNA; 967 BP.

XX ADQ30647;

XX 23-SEP-2004 (first entry)

XX West Nile virus internal diagnosis control sequence.

XX ss; internal control; West Nile Virus; diagnosis.

XX West Nile virus.

XX WO2004055159-A2.

XX 01-JUL-2004.

XX 05-DEC-2003; 2003WO-US038750.

XX 12-DEC-2002; 2002US-0432850P.

XX 20-JUN-2003; 2003US-0480431P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2004-488058/46.

XX New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.

XX Claim 27; SEQ ID NO 17; 56pp; English.

XX The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to an internal control sequence for the detection of WNV
 CC sequences using the oligonucleotides of the invention.

XX Sequence 967 BP; 273 A; 206 C; 272 G; 216 T; 0 U; 0 Other;

XX Query Match 100.0%; Score 21; DB 12; Length 967;

XX Best Local Similarity 100.0%; Pred. No. 0.65;
 XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21

Db 129 CCGGGCTGTCATATGCTAAA 149

RESULT 7

ID ADR32078 standard; DNA; 10945 BP.

XX ADR32078;

XX ADR32078;

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DT 18-NOV-2004 (first entry)
XX Genomic DNA of a West Nile virus.
XX analysis; target; real time PCR; ds; genomic.
XX West Nile virus.
XX OS
XX WO2004072230-A2.
XX PN
XX 26-AUG-2004.
XX PD
XX 10-FEB-2004; 2004WO-US002012.
XX PF
XX 10-FEB-2003; 2003US-00361004.
XX PR
XX (CLEA-) CLEARANT INC.
XX PA
XX Mckenney K, Gillmeister L, Marlowe K, Armistead D;
XX WPI; 2004-625843/60.
XX DR
XX Analyzing a target nucleic acid sequence in a biological material by real
PT time PCR using nucleic acid primers that are separated by at least 750
PT nucleic acid residues in the target sequence.
XX PS
XX Disclosure; SEQ ID NO 5; 96pp; English.
XX XX
CC The invention relates to a novel method for analysing a target nucleic
CC acid sequence in a biological material. The method comprises adding at
CC least two nucleic acid primers that hybridise under stringent conditions
CC to predetermined nucleic acid sequences of the target nucleic acid
CC sequence that are separated by at least 750 nucleic acid residues,
CC amplifying the target nucleic acid sequence by PCR, and detecting and
CC quantifying the target nucleic acid sequence. The methods and
CC compositions of the present invention are useful for analysing a target
CC nucleic acid sequence in a biological material by real time PCR using
CC nucleic acid primers that are separated by at least 750 nucleic acid
CC residues in the target sequence. This polynucleotide sequence represents
CC the genomic DNA of a West Nile virus used in the target analysis method
CC of the invention.
XX XX
SQ Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;
Query Match 100.0%; Score 21; DB 13; Length 10945;
Best Local Similarity 100.0%; Pred. No. 0.93;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CCGGGCTGTCATATGCTAAA 21
Db 87 CCGGGCTGTCATATGCTAAA 107

RESULT 8
ADR67768
ID ADR67768 standard; DNA; 10945 BP.
XX
XX ADR67768;
AC
XX
XX 18-NOV-2004 (first entry)
DT
XX
XX West Nile virus DNA detected by novel detection method.
DE
XX
XX ds; detection; pathogen.
XX
XX West Nile virus.
XX OS
XX WO2004072231-A2.
XX PN
XX 26-AUG-2004.
XX PD
XX 10-FEB-2004; 2004WO-US002013.
XX PF
XX
XX

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PR 10-FEB-2003; 2003US-00361002.
XX
XX (CLEA-) CLEARANT INC.
XX PA
XX Mckenney K, Gillmeister L, Marlowe K, Armistead D;
XX WPI; 2004-625844/60.
XX DR
XX Determining level of potentially active biological pathogens in
PT biological material, by adding nucleic acid primer pairs to biological
PT material, amplifying target nucleic acid by PCR, detecting and
PT quantifying target nucleic acid.
XX PS
XX Disclosure; SEQ ID NO 5; 11pp; English.
XX XX
CC The invention relates to a method of determining (M1) level of
CC potentially active biological pathogens in biological material, involves
CC adding at least two nucleic acid primer pairs to biological material,
CC amplifying target nucleic acid sequences by PCR, and detecting and
CC quantifying target nucleic acid sequences, where quantity of the nucleic
CC acid sequences is proportional to number of biological pathogens in
CC biological material. (M1) is useful for determining level of potentially
CC active biological pathogens in a biological material such as cells,
CC tissues, blood or blood components, proteins, enzymes, immunoglobulins,
CC botanicals, food, ligaments, tendons, nerves, bone, teeth, skin grafts,
CC bone marrow, heart valves, cartilage, corneas, arteries, veins, organs,
CC lipids, carbohydrates, collagen, chitin and its derivatives, forensic
CC samples, mummified material, human or animal remains, stem cells, islet
CC of Langerhans cells, cells for transplantation, red blood cells, white
CC blood cells or platelets. The biological pathogen is chosen from
CC bacteria, viruses, fungi and single cell parasites. The biological
CC pathogen is chosen from Aspergillus, Candida, Histoplasma,
CC Saccharomyces, Coccidioides, Cryptococcus, Escherichia, Bacillus,
CC Campylobacter, Helicobacter, Listeria, Clostridium, Streptococcus,
CC Enterococcus, Staphylococcus, Brucella, Haemophilus, Salmonella,
CC Yersinia, Pseudomonas, Serratia, Enterobacter, Klebsiella, Proteus,
CC Citrobacter, Corynebacterium, Propionibacterium and Coxiella. The
CC biological pathogen is chosen from Adeno-associated virus (AAV),
CC California encephalitis virus, Coronavirus, Coxsackievirus-A,
CC Coxsackievirus-B, Eastern equine encephalitis virus (EEEV), Echovirus,
CC Hantavirus, Hepatitis A virus (HAV), Hepatitis C virus (HCV), Hepatitis
CC delta virus (HDV), Hepatitis E virus (HEV), Hepatitis G virus (HGV), HIV,
CC Human T-lymphotrophic virus (HTLV), Influenza virus (Flu virus), Measles
CC virus (Rubella), Mumps virus, Norwalk virus, Parainfluenza virus, Polio
CC virus, Rabies virus, Respiratory Syncytial virus, Rhinovirus, Rubella
CC virus, Saint Louis encephalitis virus, Western equine encephalitis virus
CC (WEEV), Yellow fever virus, Adenovirus, Cytomegalovirus (CMV), Epstein-
CC Barr virus (EBV), Hepatitis B virus (HBV), Herpes simplex virus 1, Herpes
CC simplex virus 2, Molluscum contagiosum, Papilloma virus (HPV), Smallpox
CC virus (Variola), Vaccinia virus, Venezuelan equine encephalitis virus
CC (VEEV), Ebola virus, West Nile virus, Human Parvovirus B19 and Rotavirus.
CC (M1) is useful for determining the effectiveness of a sterilization
CC process applied to a biological material. (M1) is useful in determining
CC whether the biological pathogen is inactive or active. (M1) enables
CC determination of whether the particular biological pathogen is present in
CC a biological material as shown by amplification of first target sequence
CC and whether the biological pathogen is inactive or active. (M1) enables
CC evaluation of the effectiveness of sterilization processes, and
CC determination of both the original level and the residual level of
CC potentially active biological pathogens. This sequence corresponds to a
CC West Nile virus DNA detected by the method of the invention.
XX XX
SQ Sequence 10945 BP; 2999 A; 2457 C; 3143 G; 2346 T; 0 U; 0 Other;
Query Match 100.0%; Score 21; DB 13; Length 10945;
Best Local Similarity 100.0%; Pred. No. 0.93;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CCGGGCTGTCATATGCTAAA 21
Db 87 CCGGGCTGTCATATGCTAAA 107

```

```

RESULT 9
ADK13681
ID ADK13681 standard; DNA; 10962 BP.
XX
AC ADK13681;
XX
DT 20-MAY-2004 (first entry)
XX
DE West Nile Virus DNA sequence, SEQ ID 1.
XX
KW Virucide; Immunostimulant; flavivirus;
XX envelope protein domain III polypeptide; envelope protein; gene; ss.
XX
OS West Nile virus.
XX
FH Key Location/Qualifiers
CDS 97..10389
FT /*tag= a
FT /product= "West Nile Virus protein"
XX
PN WO2004016586-A2.
XX
PD 26-FEB-2004.
XX
PF 18-AUG-2003; 2003WO-US025681.
XX
PR 16-AUG-2002; 2002US-0403893P.
XX 06-FEB-2003; 2003US-0445581P.
XX
PA (TEXA ) UNIV TEXAS SYSTEM.
XX
PI Barrett A, Beasley D, Holbrook M;
XX
XX WPI; 2004-203756/19.
XX
DR P-PSDB; ADK13682.
XX
PT Diagnosing flavivirus infection by contacting a sample from a human or
PT animal with a flavivirus envelope protein domain III polypeptide, and
PT detecting formation of an immunocomplex between the envelope protein and
PT antibodies in the sample.
XX
PS Disclosure; SEQ ID NO 1; 110pp; English.
XX
CC The present invention relates to a method for screening for a flavivirus
CC in a subject or animal host. The method comprises: contacting a sample
CC from the subject with a composition comprising a flavivirus envelope
CC protein domain III polypeptide (ADK13683-ADK13701) under conditions that
CC permit formation of specific immunocomplex between an antibody in the
CC sample and the envelope protein domain III polypeptide; and detecting
CC whether a specific immunocomplex is formed. The present sequence is the
CC coding sequence for West Nile Virus protein, from which E protein
CC envelope protein domain III polypeptide (ADK13683) is derived.
XX
SQ Sequence 10962 BP; 2997 A; 2497 C; 3100 G; 2368 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 10962;
Best Local Similarity 100.0%; Pred. No. 0.93;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
Db 111 CCGGGCTGTCATATGCTAAA 131

RESULT 11
ABZ68481
ID ABZ68481 standard; DNA; 11029 BP.
XX
XX ABZ68481;
XX
DT 22-APR-2003 (first entry)
XX
DE Nucleotide sequence of the genome of West Nile virus IS-98-ST1.
XX
KW WNV; IS-98-ST1; Flavivirus; infection; encephalitis; gene; ss.
XX
OS West Nile virus.
XX
FH Key Location/Qualifiers
CDS 97..10397
FT /*tag= a
FT /product= "polyprotein"
XX
PN WO200281511-A1.
XX
PD 17-OCT-2002.
XX
PF 04-APR-2002; 2002WO-FR001168.
XX
ds; West Nile Virus; envelope protein; glycoprotein E; flavivirus;
Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.
West Nile virus.
WO2004040263-A2.
13-MAY-2004.
31-OCT-2003; 2003WO-US034823.
31-OCT-2002; 2002US-0422755P.
06-JUN-2003; 2003US-0476513P.
(HEAL-) HEALTH RES INC.
Wong SJ, Pei-Yong S;
WPI; 2004-400223/37.
GENBANK; AF206518.
New diagnostic kit comprising West Nile Virus (WNV) envelope protein
reactive with antibody against WNV and cross-reactive with antibody
against a flavivirus, useful in diagnosing flavivirus infection caused by
DENV, WNV, JEV or SLEV.
Disclosure; Fig 37; 212pp; English.
The invention relates to a diagnostic kit comprising at least one
isolated and purified polypeptide comprising a West Nile Virus (WNV)
envelope (E) protein or its immunogenic fragment having a native
conformation or non-denatured structure and that is reactive with
antibodies against WNV and cross-reactive with antibodies against a
flavivirus. The diagnostic kit is useful in diagnosing flavivirus
infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to
the complete nucleotide sequence of the WNV isolate 2741.
Sequence 10975 BP; 3007 A; 2460 C; 3149 G; 2359 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 10975;
Best Local Similarity 100.0%; Pred. No. 0.93;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
Db 111 CCGGGCTGTCATATGCTAAA 131

RESULT 11
ABZ68481
ID ABZ68481 standard; DNA; 11029 BP.
XX
XX ABZ68481;
XX
DT 22-APR-2003 (first entry)
XX
DE Nucleotide sequence of the genome of West Nile virus IS-98-ST1.
XX
KW WNV; IS-98-ST1; Flavivirus; infection; encephalitis; gene; ss.
XX
OS West Nile virus.
XX
FH Key Location/Qualifiers
CDS 97..10397
FT /*tag= a
FT /product= "polyprotein"
XX
PN WO200281511-A1.
XX
PD 17-OCT-2002.
XX
PF 04-APR-2002; 2002WO-FR001168.
XX
West Nile Virus isolate 2741 complete genome sequence.

```


XX 04-APR-2001; 2001FR-00004599.
 PR 06-SEP-2001; 2001FR-00011525.
 XX (INSP) INST PASTEUR.
 PA (KIMR-) KIMRON VETERINARY INST.
 XX Despres P, Deubel V, Guenet J, Drouet M, Malkinson M, Banet C;
 PI Frenkiel M, Courageot M, Coulibaly F, Catteau A, Flamand M, Weber P;
 PI Ceccaldi P;
 XX WPI; 2003-058498/05.
 DR P-PSDB; ABP70647.
 DR New neurovirulent strain of West Nile virus, useful in diagnosis and
 XX screening for antiviral agents, also related nucleic acids, proteins and
 PT antibodies.
 XX Claim 1; Page 34-49; 68pp; French.
 PS The present sequence represents the genome of a strain of West Nile virus
 XX (WNV), designated IS-98-ST1. This strain is a neuroinvasive and
 CC neurovirulent strain of WNV. Polynucleotides and polypeptides derived
 CC from the IS-98-ST1 genome are useful for diagnosis and prognosis of
 CC Flavivirus infection, specifically WNV-mediated encephalitis. They are
 CC also useful to raise specific antibodies, for recombinant expression of
 CC WNV proteins or peptides (for diagnosis, production of antibodies and
 CC identification of specific binding partners in cells), for identifying
 CC cellular genes implicated in resistance to viral infection, and for
 CC screening for anti-Flavivirus agents
 XX Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 21; DB 8; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 0.93;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CCGGGCTGTCATATGCTAAA 21
 Db 129 CCGGGCTGTCATATGCTAAA 149
 RESULT 12
 ABV74821
 ID ABV74821 standard; DNA; 11029 BP.
 XX AC ABV74821;
 XX 28-MAR-2003 (first entry)
 DT West Nile virus strain NY99-flamingo 382-99 complete genome.
 DE Virucide; hepatotropic; antiinflammatory; antiviral; OAS;
 XX 2'-5'-oligoadenylate synthase; Flavivirus infection; gene; ss.
 KW West Nile Virus.
 OS Key Location/Qualifiers
 XX CDS 97..10398
 FT /*tag= a
 FT /product= "West Nile Virus protein"
 XX WO200281741-A2.
 XX 17-OCT-2002.
 PD 04-APR-2002; 2002WO-FR001169.
 XX 04-APR-2001; 2001FR-00004598.
 XX (INSP) INST PASTEUR.
 PA (CNRS) CNRS CENT NAT RECH SCI.
 XX

PI Guenet J, Mashimo T, Simon-Chazottes D, Montagutelli X;
 PI Frenkiel M, Despres P, Deubel V, Bonhomme F, Lucas M;
 XX WPI; 2003-058566/05.
 DR P-PSDB; ABB98821.
 XX Identifying stimulators of oligoadenylate synthase family genes, useful
 PT as antiviral agents against Flavivirus, also mutated genes responsible
 PT for sensitivity to virus.
 XX Example 1; Page 52-67; 93pp; French.
 PS The present invention relates to a method for identifying compounds (I)
 CC that can stimulate a gene of the OAS (2'-5'-oligoadenylate synthase)
 CC family. The method comprises: (a) inducing expression of the OAS gene in
 CC a culture of cells from a non-human mammal (Flvr/Flvr or Flvr/Flvs;
 CC indicating resistance or sensitivity to Flavivirus infection); (b)
 CC treating cells with test compound; and (c) measuring activity of OAS gene
 CC relative to a control. (I) are potentially useful as antiviral agents for
 CC treating infections by Flaviviruses (e.g. hepatitis C; dengue; yellow
 CC fever and various forms of encephalitis). Genomic OAS DNA and derived
 CC cDNA, also the encoded proteins, are useful: (a) for treating Flavivirus
 CC infection; (b) in screening for anti-Flavivirus agents, and (c) for
 CC evaluating sensitivity of subjects to Flavivirus infection and their
 CC likely response to interferon treatment, e.g. to identify patients at
 CC risk of developing severe forms of such infections. The present sequence
 CC is West Nile Virus strain NY99-flamingo 382-99 (IS-98-ST1) complete
 CC genome, which was used in an example from the invention. West Nile Virus
 CC is one such Flavivirus
 XX Sequence 11029 BP; 3019 A; 2471 C; 3167 G; 2372 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 21; DB 10; Length 11029;
 Best Local Similarity 100.0%; Pred. No. 0.93;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CCGGGCTGTCATATGCTAAA 21
 Db 129 CCGGGCTGTCATATGCTAAA 149
 RESULT 13
 ADN98023
 ID ADN98023 standard; DNA; 11029 BP.
 XX AC ADN98023;
 XX 29-JUL-2004 (first entry)
 DT West Nile Virus isolate 3356 complete genome sequence.
 DE ds; West Nile Virus; envelope protein; glycoprotein E; flavivirus;
 XX Japanese encephalitis virus; Dengue virus; St Louis encephalitis virus.
 KW West Nile virus.
 OS WO2004040263-A2.
 XX 13-MAY-2004.
 XX 31-OCT-2003; 2003WO-US034823.
 PF 31-OCT-2002; 2002US-0422755P.
 PR 06-JUN-2003; 2003US-0476513P.
 XX (HEAL-) HEALTH RES INC.
 XX Wong SJ, Pei-Yong S;
 PI WPI; 2004-400223/37.
 DR GENBANK; AF04756.
 XX New diagnostic kit comprising West Nile Virus (WNV) envelope protein

PT reactive with antibody against WNV and cross-reactive with antibody
PT against a flavivirus, useful in diagnosing flavivirus infection caused by
PT DENV, WNV, JEV or SLEV.

XX Disclosure; Fig 38; 212pp; English.

XX The invention relates to a diagnostic kit comprising at least one
CC isolated and purified polypeptide comprising a West Nile Virus (WNV)
CC envelope (E) protein or its immunogenic fragment having a native
CC conformation or non-denatured structure and that is reactive with a
CC antibodies against WNV and cross-reactive with antibodies against a
CC flavivirus. The diagnostic kit is useful in diagnosing flavivirus
CC infection caused by DENV, WNV, JEV or SLEV. This sequence corresponds to
XX the complete nucleotide sequence of the WNV isolate 3356.

XX Sequence 11029 BP; 3017 A; 2466 C; 3172 G; 2374 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 12; Length 11029;
Best Local Similarity 100.0%; Pred. No. 0.93;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCGGGCTGTCATATGCTAAA 21
Db 129 CCGGGCTGTCATATGCTAAA 149

RESULT 14
ADN36695/c
ID ADN36695 standard; DNA; 21 BP.
XX
XX AC ADN36695;
XX
XX 15-JUL-2004 (first entry)
XX
XX West Nile virus detection-related oligonucleotide probe SeqID17.
XX
XX hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.

XX West Nile virus.

XX WO2004036190-A2.

XX 29-APR-2004.

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

XX 25-NOV-2002; 2002US-0429006P.

XX 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Disclosure; SEQ ID NO 17; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental

CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.

XX Sequence 21 BP; 5 A; 5 C; 4 G; 7 T; 0 U; 0 Other;

Query Match 90.5%; Score 19; DB 12; Length 21;
Best Local Similarity 100.0%; Pred. No. 4.4;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GGGCTGTCATATGCTAAA 21
Db 21 GGGCTGTCATATGCTAAA 3

RESULT 15
ADN36823/c
ID ADN36823 standard; RNA; 24 BP.

XX
XX AC ADN36823;

XX 15-JUL-2004 (first entry)

XX West Nile virus detection-related oligonucleotide probe SeqID145.

XX hybridisation assay probe; nucleic acid detection;
KW target-complementary sequence; flavivirus; West Nile virus; WNV;
KW RNA virus; infection; meningitis; encephalitis;
KW high throughput screening; probe; ss.

XX West Nile virus.

XX Key Location/Qualifiers

XX modified_base 1..24

XX /*tag= a

XX /mod_base= OTHER

XX /note= "OTHER= 2'-methoxyethoxy (2'-MOE) nucleotides"

XX WO2004036190-A2.

XX 29-APR-2004.

XX 10-OCT-2003; 2003WO-US033639.

XX 16-OCT-2002; 2002US-0418891P.

XX 25-NOV-2002; 2002US-0429006P.

XX 24-FEB-2003; 2003US-0449810P.

XX (GENP-) GEN-PROBE INC.

XX Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;

XX WPI; 2004-389590/36.

XX New hybridization assay probe comprising target-complementary sequence of
PT bases, useful in detecting flavivirus, e.g. West Nile virus.

XX Example 1; SEQ ID NO 145; 135pp; English.

XX This invention relates to a novel hybridisation assay probe, for
CC detecting a nucleic acid, which is a probe sequence that comprises a
CC target-complementary sequence of bases, and optionally one or more base
CC sequences that are not complementary to the nucleic acid that is to be
CC detected. The hybridisation assay probes and the kits are useful in
CC detecting and amplifying a target nucleic acid sequence, for example
CC flavivirus like West Nile virus, that may be present in a biological
CC sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC birds and culex mosquitoes, with humans and horses serving as incidental
CC hosts. Infection of humans can lead to meningitis or encephalitis. The
CC invention may allow for accurate and efficient high throughput screening.
CC The present sequence is that of an oligonucleotide probe which is related
CC to the invention.

```

XX SQ Sequence 24 BP; 5 A; 7 C; 5 G; 0 T; 7 U; 0 Other;
Query Match 90.5%; Score 19; DB 12; Length 24;
Best Local Similarity 100.0%; Pred. No. 4.4;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 GGGCTGTCAATATGCTAAA 21
Db 24 GGGCTGTCAATATGCTAAA 6

RESULT 16
ADN36707/c
ID ADN36707 standard; DNA; 48 BP.
XX AC
XX AC ADN36707;
XX DT
XX DT 15-JUL-2004 (first entry)
XX DE
XX DE West Nile virus detection-related oligonucleotide probe SeqID29.
XX KW hybridisation assay probe; nucleic acid detection;
XX KW target-complementary sequence; flavivirus; West Nile virus; WNV;
XX KW RNA virus; infection; meningitis; encephalitis;
XX KW high throughput screening; probe; ss.
XX OS
XX OS West Nile virus.
XX OS Enterobacteria phage T7.
XX FH
XX FH Key Location/Qualifiers
FT misc_feature 1..27
FT /*tag= a
FT /note= "T7 promoter sequence"
FT misc_feature 28..48
FT /*tag= b
FT /note= "WNV-complementary sequence"
XX FT
XX FT
XX W02004036190-A2.
XX PN
XX PD
XX PD 29-APR-2004.
XX PF
XX PF 10-OCT-2003; 2003WO-US033639.
XX PR
XX PR 16-OCT-2002; 2002US-0418891P.
XX PR 25-NOV-2002; 2002US-0429006P.
XX PR 24-FEB-2003; 2003US-0449810P.
XX XX
XX XX (GENP-) GEN-PROBE INC.
XX PI
XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX XX
XX XX New hybridization assay probe comprising target-complementary sequence of
XX bases, useful in detecting flavivirus, e.g. West Nile virus.
XX PS
XX PS Disclosure; SEQ ID NO 29; 135pp; English.
XX XX
XX XX This invention relates to a novel hybridisation assay probe, for
XX detecting a nucleic acid, which is a probe sequence that comprises a
XX target-complementary sequence of bases, and optionally one or more base
XX sequences that are not complementary to the nucleic acid that is to be
XX detected. The hybridisation assay probes and the kits are useful in
XX detecting and amplifying a target nucleic acid sequence, for example
XX flavivirus like West Nile virus, that may be present in a biological
XX sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX birds and culex mosquitoes, with humans and horses serving as incidental
XX hosts. Infection of humans can lead to meningitis or encephalitis. The
XX invention may allow for accurate and efficient high throughput screening.
XX The present sequence is that of an oligonucleotide probe which is related
XX to the invention.

SQ Sequence 48 BP; 16 A; 9 C; 9 G; 14 T; 0 U; 0 Other;
Query Match 90.5%; Score 19; DB 12; Length 48;
Best Local Similarity 100.0%; Pred. No. 4.9;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 GGGCTGTCAATATGCTAAA 21
Db 48 GGGCTGTCAATATGCTAAA 30

RESULT 17
ADN36694/c
ID ADN36694 standard; DNA; 69 BP.
XX AC
XX AC ADN36694;
XX DT
XX DT 15-JUL-2004 (first entry)
XX DE
XX DE West Nile virus detection-related oligonucleotide probe SeqID16.
XX KW hybridisation assay probe; nucleic acid detection;
XX KW target-complementary sequence; flavivirus; West Nile virus; WNV;
XX KW RNA virus; infection; meningitis; encephalitis;
XX KW high throughput screening; probe; ss.
XX OS
XX OS West Nile virus.
XX PN
XX PN W02004036190-A2.
XX PD
XX PD 29-APR-2004.
XX PF
XX PF 10-OCT-2003; 2003WO-US033639.
XX PR
XX PR 16-OCT-2002; 2002US-0418891P.
XX PR 25-NOV-2002; 2002US-0429006P.
XX PR 24-FEB-2003; 2003US-0449810P.
XX XX
XX XX (GENP-) GEN-PROBE INC.
XX PI
XX PI Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
XX WPI; 2004-389590/36.
XX XX
XX XX New hybridization assay probe comprising target-complementary sequence of
XX bases, useful in detecting flavivirus, e.g. West Nile virus.
XX PS
XX PS Claim 68; SEQ ID NO 16; 135pp; English.
XX XX
XX XX This invention relates to a novel hybridisation assay probe, for
XX detecting a nucleic acid, which is a probe sequence that comprises a
XX target-complementary sequence of bases, and optionally one or more base
XX sequences that are not complementary to the nucleic acid that is to be
XX detected. The hybridisation assay probes and the kits are useful in
XX detecting and amplifying a target nucleic acid sequence, for example
XX flavivirus like West Nile virus, that may be present in a biological
XX sample. West Nile virus (WNV) is an RNA virus that primarily infects
XX birds and culex mosquitoes, with humans and horses serving as incidental
XX hosts. Infection of humans can lead to meningitis or encephalitis. The
XX invention may allow for accurate and efficient high throughput screening.
XX The present sequence is that of an oligonucleotide probe which is related
XX to the invention.

SQ Sequence 69 BP; 18 A; 22 C; 14 G; 15 T; 0 U; 0 Other;
Query Match 90.5%; Score 19; DB 12; Length 69;
Best Local Similarity 100.0%; Pred. No. 5.2;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 GGGCTGTCAATATGCTAAA 21
Db 69 GGGCTGTCAATATGCTAAA 51

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RESULT 18
ADN36696/c
ID  ADN36696 standard; DNA; 20 BP.
XX
XX  ADN36696;
AC
XX
XX  15-JUL-2004 (first entry)
DT
XX
XX  West Nile virus detection-related oligonucleotide probe SeqID18.
DE
XX
XX  hybridisation assay probe; nucleic acid detection;
KW  target-complementary sequence; flavivirus; West Nile virus; WNV;
KW  RNA virus; infection; meningitis; encephalitis;
KW  high throughput screening; probe; ss.
XX
XX  West Nile virus.
OS
XX
XX  WO2004036190-A2.
PN
XX
XX  29-APR-2004.
PD
XX
XX  10-OCT-2003; 2003WO-US033639.
PF
XX
XX  16-OCT-2002; 2002US-0418891P.
PR
XX
XX  25-NOV-2002; 2002US-0429006P.
PR
XX
XX  24-FEB-2003; 2003US-0449810P.
PR
XX
XX  (GENP-) GEN-PROBE INC.
PA
XX
XX  Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
PI
XX
XX  WPI; 2004-389590/36.
DR
XX
XX  New hybridization assay probe comprising target-complementary sequence of
PT  bases, useful in detecting flavivirus, e.g. West Nile virus.
PT
XX
XX  Disclosure; SEQ ID NO 18; 135pp; English.
PS
XX
XX  This invention relates to a novel hybridisation assay probe, for
CC  detecting a nucleic acid, which is a probe sequence that comprises a
CC  target-complementary sequence of bases, and optionally one or more base
CC  sequences that are not complementary to the nucleic acid that is to be
CC  detected. The hybridisation assay probes and the kits are useful in
CC  detecting and amplifying a target nucleic acid sequence, for example
CC  flavivirus like West Nile virus, that may be present in a biological
CC  sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC  birds and culex mosquitoes, with humans and horses serving as incidental
CC  hosts. Infection of humans can lead to meningitis or encephalitis. The
CC  invention may allow for accurate and efficient high throughput screening.
CC  The present sequence is that of an oligonucleotide probe which is related
CC  to the invention.
CC
XX  Query Match 85.7%; Score 18; DB 12; Length 20;
SQ  Sequence 20 BP; 5 A; 4 C; 4 G; 7 T; 0 U; 0 Other;
    Best Local Similarity 100.0%; Pred. No. 15;
    Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
    Qy 4 GGCTGTCATATGCTAAA 21
    Db 20 GGCTGTCATATGCTAAA 3
    RESULT 19
    ADN36708/c
    ID  ADN36708 standard; DNA; 47 BP.
    XX
    XX  ADN36708;
    AC
    XX
    XX  15-JUL-2004 (first entry)
    DT
    XX
    XX  West Nile virus detection-related oligonucleotide probe SeqID30.
    DE
```

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XX  hybridisation assay probe; nucleic acid detection;
KW  target-complementary sequence; flavivirus; West Nile virus; WNV;
KW  RNA virus; infection; meningitis; encephalitis;
KW  high throughput screening; probe; ss.
XX
XX  West Nile virus.
OS
XX
XX  Enterobacteria phage T7.
XX
XX  Key Location/Qualifiers
FT  misc_feature 1..27
FT  /*tag= a
FT  /note= "T7 promoter sequence"
FT  misc_feature 28..47
FT  /*tag= b
FT  /note= "WNV-complimentary sequence"
XX
XX  WO2004036190-A2.
PN
XX
XX  29-APR-2004.
PD
XX
XX  10-OCT-2003; 2003WO-US033639.
PF
XX
XX  16-OCT-2002; 2002US-0418891P.
PR
XX
XX  25-NOV-2002; 2002US-0429006P.
PR
XX
XX  24-FEB-2003; 2003US-0449810P.
PR
XX
XX  (GENP-) GEN-PROBE INC.
PA
XX
XX  Linnen JM, Pollner RB, Wu W, Dennis GG, Darby PM;
PI
XX
XX  WPI; 2004-389590/36.
DR
XX
XX  New hybridization assay probe comprising target-complementary sequence of
PT  bases, useful in detecting flavivirus, e.g. West Nile virus.
PT
XX
XX  Disclosure; SEQ ID NO 30; 135pp; English.
PS
XX
XX  This invention relates to a novel hybridisation assay probe, for
CC  detecting a nucleic acid, which is a probe sequence that comprises a
CC  target-complementary sequence of bases, and optionally one or more base
CC  sequences that are not complementary to the nucleic acid that is to be
CC  detected. The hybridisation assay probes and the kits are useful in
CC  detecting and amplifying a target nucleic acid sequence, for example
CC  flavivirus like West Nile virus, that may be present in a biological
CC  sample. West Nile virus (WNV) is an RNA virus that primarily infects
CC  birds and culex mosquitoes, with humans and horses serving as incidental
CC  hosts. Infection of humans can lead to meningitis or encephalitis. The
CC  invention may allow for accurate and efficient high throughput screening.
CC  The present sequence is that of an oligonucleotide probe which is related
CC  to the invention.
CC
XX  Query Match 85.7%; Score 18; DB 12; Length 47;
SQ  Sequence 47 BP; 16 A; 8 C; 9 G; 14 T; 0 U; 0 Other;
    Best Local Similarity 100.0%; Pred. No. 17;
    Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
    Qy 4 GGCTGTCATATGCTAAA 21
    Db 47 GGCTGTCATATGCTAAA 30
    RESULT 20
    AAQ22767
    ID  AAQ22767 standard; DNA; 4512 BP.
    XX
    XX  AAQ22767;
    AC
    XX
    XX  12-AUG-1992 (first entry)
    DT
    XX
    XX  JEV Nakayama strain prW, E, NS1, NS2A, NS2B and C coding regions.
    XX
```

KW Japanese Encephalitis Virus; vaccinia virus donor; plasmid pDr20; ss.
 XX OS
 XX Japanese encephalitis virus.
 XX PN WO9203545-A.
 XX PD 05-MAR-1992.
 XX PF 05-AUG-1991; 91WO-US005816.
 XX PR 15-AUG-1990; 90US-00567960.
 XX PR 06-JUN-1991; 91US-00711429.
 XX PR 13-JUN-1991; 91US-00714687.
 XX PR 17-JUL-1991; 91US-00729800.
 XX PR 05-AUG-1991; 91WO-U0005816.
 XX PA (VIRO-) VIROGENETICS CORP.
 XX PI Paoletti E, Pinc, Pinc, Pincus SE;
 XX PS WPI; 1992-096889/12.
 XX PT Recombinant pox-virus e.g. vaccinia, fowl-pox and canary-pox virus -
 PT contg. DNA from flavi-virus e.g. Japanese encephalitis and yellow fever
 PT virus, used as vaccine.
 XX PS Example 9; Fig 17; 117pp; English.
 XX CC CDNA was prepared from genomic virion RNA obtained from suspension
 CC cultures of C6/36 cells infected with a passage 55 suckling mouse brain
 CC stock of the Nagayama strain of JEV. EcoRI linkers were ligated to the
 CC cDNA fragments for cloning into pBR322. Recombinant plasmids were
 CC transformed into E.coli DH5 cells. Plasmid pC20 contained 81 non-coding
 CC nucleotides and the coding regions for C and pM. Sequence AAQ22767 is
 CC that of the C coding region of pC20, combined with an updated sequence of
 CC the pM, E, NS1, NS2A and NS2B coding regions of the Nagayama strain. The
 CC sequence begins at the C protein Met initiation codon. A subfragment of
 CC pC20 was cloned into pUC18 to give pDr20. This plasmid was then used in
 CC the construction of novel recombinants JEV24, JEV27, JEV33 and JEV34.
 CC These were transfected into VP410 infected cells to generate VP825,
 CC VP829, VP857 and VP864, respectively
 XX SQ Sequence 4512 BP; 1192 A; 1055 C; 1253 G; 1012 T; 0 U; 0 Other;
 Query Match 84.8%; Score 17.8; DB 2; Length 4512;
 Best Local Similarity 90.5%; Pred. No. 41;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 CCGGGCTGCTCAATGCTGCTAAA 21
 Db 33 CCGGGCTGCTCAATGCTGCTAAA 53
 RESULT 21
 ADO07431
 ID ADO07431 standard; DNA; 10818 BP.
 XX AC ADO07431;
 XX DT 15-JUL-2004 (first entry)
 XX DE Japanese Encephalitis virus JEV coding sequence SEQ ID NO: 9.
 XX DE antiinflammatory; neuroprotective; gene therapy;
 KW Japanese Encephalitis virus; JEV; ds; gene; vaccine;
 KW Japanese encephalitis.
 XX OS Japanese encephalitis virus.
 XX PN WO2004033690-A1.
 XX PD 22-APR-2004.

PF 09-OCT-2003; 2003WO-KR002081.
 XX PR 09-OCT-2002; 2002KR-00061589.
 XX PA (CDC-) CID CO LTD.
 XX PA (LEES/) LEE S H.
 XX PI Lee SH, Lee Y, Yun S;
 XX DR WPI; 2004-340933/31.
 XX PT New Japanese encephalitis virus genomic RNA, useful in developing
 PT vaccines for and in diagnosing and treating Japanese encephalitis.
 XX PS Example 2; Page 145-152; 265pp; English.
 XX CC The present invention relates to a genomic RNA of the Korean Japanese
 CC Encephalitis Virus (JEV) isolate, composed of a 5' and 3' nontranslated
 CC region (NTR) and a single polypeptide coding region. The JEV genomic RNA,
 CC JEV cDNA and reagents are useful in developing vaccines for and in
 CC diagnosing and treating Japanese encephalitis. The present sequence is a
 CC sequence of the invention.
 XX SQ Sequence 10818 BP; 2991 A; 2491 C; 3075 G; 2261 T; 0 U; 0 Other;
 Query Match 84.8%; Score 17.8; DB 12; Length 10818;
 Best Local Similarity 90.5%; Pred. No. 47;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 CCGGGCTGCTCAATGCTGCTAAA 21
 Db 78 CCGGGCTGCTCAATGCTGCTAAA 98
 RESULT 22
 ADO07437
 ID ADO07437 standard; DNA; 10968 BP.
 XX AC ADO07437;
 XX DT 15-JUL-2004 (first entry)
 XX DE Japanese Encephalitis virus JEV coding sequence SEQ ID NO: 15.
 XX DE antiinflammatory; neuroprotective; gene therapy;
 KW Japanese Encephalitis virus; JEV; ds; gene; vaccine;
 KW Japanese encephalitis.
 XX OS Japanese encephalitis virus.
 XX PN WO2004033690-A1.
 XX PD 22-APR-2004.
 XX PF 09-OCT-2003; 2003WO-KR002081.
 XX PR 09-OCT-2002; 2002KR-00061589.
 XX PA (CDC-) CID CO LTD.
 XX PA (LEES/) LEE S H.
 XX PI Lee SH, Lee Y, Yun S;
 XX DR WPI; 2004-340933/31.
 XX PT New Japanese encephalitis virus genomic RNA, useful in developing
 PT vaccines for and in diagnosing and treating Japanese encephalitis.
 XX PS Claim 3; Page 154-161; 265pp; English.
 XX CC The present invention relates to a genomic RNA of the Korean Japanese
 CC Encephalitis Virus (JEV) isolate, composed of a 5' and 3' nontranslated
 CC region (NTR) and a single polypeptide coding region. The JEV genomic RNA,


```

Qy 1 CCGGCTGTCATATGCTAAA 21
    |||||
Db 128 CCGGCTATCAATATGCTGAA 148

```

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 17:45:55 ; Search time 1500.84 Seconds
(without alignments)
532.600 Million cell updates/sec

Title: US-10-729-421-34
Perfect score: 21
Sequence: 1 ccgggctgctcaatgctaaa 21

Scoring table: IDENTITY NUC
Gapop 10'0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 100 summaries

Database : EST:*

- 1: gb_est1:*
- 2: gb_est2:*
- 3: gb_hic:*
- 4: gb_est3:*
- 5: gb_est4:*
- 6: gb_est5:*
- 7: gb_est6:*
- 8: gb_gse1:*
- 9: gb_gse2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|--------------------|
| 1 | 17.8 | 84.8 | 425 | 7 | C0951450 UMC-pd12f |
| 2 | 17.8 | 84.8 | 696 | 7 | CN153647 940784 MA |
| 3 | 17.8 | 84.8 | 696 | 7 | CN155761 943088 MA |
| 4 | 17.8 | 84.8 | 998 | 9 | AG570358 Mus muscu |
| 5 | 17.4 | 82.9 | 429 | 7 | CK699541 ZF101-P00 |
| 6 | 17.4 | 82.9 | 693 | 9 | CB434484 tigr-ges- |
| 7 | 17 | 81.0 | 1009 | 9 | CN808327 Drosophil |
| 8 | 16.8 | 80.0 | 107 | 9 | CK091023 F039P30.3 |
| 9 | 16.8 | 80.0 | 354 | 7 | CK091023 F039P30.3 |
| 10 | 16.8 | 80.0 | 367 | 7 | CK101326 F039P30.5 |
| 11 | 16.8 | 80.0 | 407 | 4 | BG037611 dcs3e09.Y |
| 12 | 16.8 | 80.0 | 501 | 7 | CN489600 Mdfw20191 |
| 13 | 16.8 | 80.0 | 516 | 8 | AZ981379 2M0258113 |
| 14 | 16.8 | 80.0 | 562 | 5 | BUB15985 N058F04 P |
| 15 | 16.8 | 80.0 | 620 | 6 | CA821839 RSH08C06 |
| 16 | 16.8 | 80.0 | 623 | 8 | AZ832056 2M0112L09 |
| 17 | 16.8 | 80.0 | 631 | 7 | CK317796 B9F01N01 |
| 18 | 16.8 | 80.0 | 635 | 5 | BUB63468 S028D11 P |
| 19 | 16.8 | 80.0 | 751 | 7 | CV257160 WS0245.B2 |
| 20 | 16.8 | 80.0 | 790 | 9 | CR268537 Reverse a |
| 21 | 16.8 | 80.0 | 900 | 9 | CR016265 Forward a |
| 22 | 16.8 | 80.0 | 986 | 5 | BQ220271 AGENCOURT |
| 23 | 16.4 | 78.1 | 298 | 9 | CE199293 tigr-ges- |
| 24 | 16.4 | 78.1 | 363 | 1 | AL926049 AL926049 |

| | | | | |
|-----------|------------|------|---|-----------|
| AI512957 | LD45125.5 | 502 | 1 | AI512957 |
| BX723363 | BX723363 | 522 | 5 | BX723363 |
| BH039924 | RPCI-24-2 | 604 | 8 | BH039924 |
| CO125121 | GR_Eb08G | 686 | 7 | CO125121 |
| CC870522 | NDL_123C7 | 704 | 9 | CC870522 |
| BU381267 | 603861468 | 815 | 5 | BU381267 |
| BX723938 | BX723938 | 851 | 5 | BX723938 |
| BI320339 | sa21f09. | 223 | 4 | BI320339 |
| CF190539 | k8k07j2.f | 403 | 7 | CF190539 |
| CN496900 | Mdfw20230 | 405 | 7 | CN496900 |
| CO285402 | EK168918. | 432 | 7 | CO285402 |
| AA918584 | OL53a09.a | 503 | 1 | AA918584 |
| AZ713066 | RPCI-24-6 | 537 | 8 | AZ713066 |
| BY566461 | BY566461 | 543 | 6 | BY566461 |
| AQ425764 | CITBI-EI- | 551 | 8 | AQ425764 |
| CE166745 | tigr-ges- | 560 | 9 | CE166745 |
| BX711144 | BX711144 | 570 | 5 | BX711144 |
| CN995009 | Mdfw2050m | 573 | 7 | CN995009 |
| BF260707 | HVSNEF002 | 577 | 2 | BF260707 |
| BG000437 | MR3--GN015 | 578 | 4 | BG000437 |
| BM440929 | EBro08_SQ | 592 | 4 | BM440929 |
| BF614355 | de05e10.Y | 601 | 2 | BF614355 |
| BF676682 | 602086323 | 618 | 2 | BF676682 |
| CA694474 | wlnk4_pk0 | 621 | 6 | CA694474 |
| CN910418 | 030128ABL | 637 | 7 | CN910418 |
| BJS589589 | BJS589589 | 651 | 4 | BJS589589 |
| CF365321 | 835940_MA | 652 | 7 | CF365321 |
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| BJS589606 | BJS589606 | 698 | 4 | BJS589606 |
| AG417300 | Mus muscu | 721 | 9 | AG417300 |
| AG558847 | Mus muscu | 732 | 9 | AG558847 |
| AG548062 | Mus muscu | 743 | 9 | AG548062 |
| AG566555 | Mus muscu | 744 | 9 | AG566555 |
| BJS594699 | BJS594699 | 756 | 4 | BJS594699 |
| CO802419 | AGENCOURT | 757 | 7 | CO802419 |
| BX997985 | Reverse a | 800 | 9 | BX997985 |
| AZ187754 | SP_1009_B | 824 | 8 | AZ187754 |
| BZ701219 | PUCO20TD | 859 | 8 | BZ701219 |
| BZ704489 | PUCO20TD | 860 | 8 | BZ704489 |
| CF374332 | AGENCOURT | 863 | 7 | CF374332 |
| BX851663 | BX851663 | 865 | 5 | BX851663 |
| CL240184 | ZMWBb058 | 932 | 9 | CL240184 |
| BQ688620 | AGENCOURT | 936 | 5 | BQ688620 |
| CNS04CON | Tetraodon | 939 | 9 | CNS04CON |
| CD245530 | AGENCOURT | 942 | 6 | CD245530 |
| CG094980 | PUKBV25TD | 953 | 9 | CG094980 |
| CG094979 | PUKBV26TB | 1009 | 9 | CG094979 |
| CL997901 | ZMWBHf001 | 1016 | 9 | CL997901 |
| CC221736 | CH261-921 | 1049 | 9 | CC221736 |
| CC221316 | CH261-83N | 1058 | 8 | CC221316 |
| BI028233 | CM4-MT028 | 1062 | 8 | BI028233 |
| CB218410 | NISC_nb08 | 391 | 4 | CB218410 |
| AQ226974 | HS_2013_A | 402 | 6 | AQ226974 |
| CB471568 | tigr-ges- | 535 | 8 | CB471568 |
| BH352121 | CH230-220 | 657 | 8 | BH352121 |
| CK448276 | pncs900ad | 774 | 7 | CK448276 |
| CL292196 | Tetraodon | 929 | 9 | CL292196 |
| CL477040 | SAIL_266 | 959 | 9 | CL477040 |
| CL046976 | CH216-65F | 1050 | 9 | CL046976 |
| CD112634 | MEI-0024T | 220 | 6 | CD112634 |
| R40815 | Yf82b01.81 | 234 | 7 | R40815 |
| R61488 | Yf15g09.81 | 246 | 7 | R61488 |
| BP751208 | BP751208 | 268 | 5 | BP751208 |
| AV210948 | AV210948 | 270 | 1 | AV210948 |
| HO9942 | ym01c11.81 | 273 | 7 | HO9942 |
| AA958331 | MAAD0514. | 286 | 1 | AA958331 |
| AV228707 | AV228707 | 294 | 1 | AV228707 |
| CC072824 | CSU_K33r. | 295 | 8 | CC072824 |
| B1126863 | I081P66P | 305 | 4 | B1126863 |
| B1126556 | I077P15P | 328 | 4 | B1126556 |
| AZ783049 | 2M0024J20 | 340 | 8 | AZ783049 |
| AU233672 | AU233672 | 350 | 1 | AU233672 |

98 15.8 75.2 355 1 AU278073 AU278073
 c 99 15.8 75.2 388 9 CE627437 tigr-gss-
 c 100 15.8 75.2 430 8 AQ437513 HS_5133_B

ALIGNMENTS

RESULT 1
 CO951450 425 bp mRNA linear EST 09-SEP-2004
 LOCUS UMC-pd12fol-008-a09 Day 12 ovarian follicle pd12fol Sus scrofa cDNA
 DEFINITION 3', mRNA sequence.
 ACCESSION CO951450
 VERSION CO951450.1 GI:51328453
 KEYWORDS EST.
 SOURCE Sus scrofa (pig)
 ORGANISM Sus scrofa
 Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
 1 (bases 1 to 425)
 Jiang,H., Whitworth,K.M., Bivens,N.J., Ries,J.E., Woods,R.J.,
 Forrester,L.J., Springer,G.K., Mathialagan,N., Agca,C.,
 Prather,R.S. and Lucy,M.C.
 Large-scale Generation and Analysis of Expressed Sequence Tags from
 Porcine Ovary
 Unpublished (2004)
 Contact: DNA Core Facility (Swine Project)
 Animal Science - RS Prather
 University of Missouri-Columbia
 M616 Medical Sciences Bldg., Columbia, MO 65212, USA
 Tel: (573)882-0428
 Fax: (573)884-5552
 Email: porcine@rnet.missouri.edu
 POLYA=No.

FEATURES

Location/Qualifiers
 1..425
 /organism="Sus scrofa"
 /mol_type="mRNA"
 /db_xref="taxon:9823"
 /dev_stages="Day 12 ovarian follicle"
 /clone_libs="pd12fol"
 /note="Vector: T37PAC; Funding: A grant from the Monsanto Company to the University of Missouri. Genetic Source: Ovarian tissue (whole ovary, dissected follicles, or corpora lutea) was collected from crossbred pigs (Sus scrofa domestica), frozen in liquid nitrogen shortly after collection, and stored at -80 degrees Celsius until RNA extraction. The tissue from several individual pigs was pooled for the purpose of RNA extraction. The specific tissues collected were fetal whole ovary; neonatal whole ovary; prepubertal whole ovary; 2, 4, 6 and 8 mm growing follicles; Day 0 follicles; Day 5 small antral follicles and corpora lutea; Day 12 corpora lutea and Day 12 follicles. More information regarding the methods can be found at:
<http://genome.rnet.missouri.edu/Swine/Methods.html>.
 Library Construction (Standard Protocol): All procedures discussed in this section have been described in detail elsewhere (Soares et al., 1994; Bonaldo et al., 1996; Jiang et al., 2001). Total cellular RNA from each sample was isolated by using STAT-60 reagent (Tel-Test, Friendswood, TX) and the poly(A)+ RNA was obtained by two rounds of purification with the Oligotex mRNA isolation kit (Qiagen) according to the manufacturer's instructions. The libraries were constructed essentially as described by the manufacturer's instructions provided with the SuperScript Plasmid System (Invitrogen, cat. no. 18248-013). Briefly, 1mg of poly(A)+ RNA will be annealed at 37 degrees Celsius with 10mg of NotI-tag-dT18 oligonucleotide (GCTGCTCGGCCGC-tag-r18) and reverse transcribed at 37 degrees Celsius with SuperScript II (Invitrogen) reverse transcriptase (Jiang et al., 2001).

The 'tag' represents a tissue/stage-specific ten-base sequence identifier (<http://genome.uiowa.edu/pubsoft/software.html>) present in the oligonucleotide used to prime first-strand synthesis. Second strand synthesis was performed with T4 DNA polymerase in the presence of DNA ligase and RNase H. After second strand synthesis, the double-stranded cDNAs was ligated to SalI adapters (Invitrogen-Life Technologies) and digested with NotI. The cDNAs will be size selected by passage through cDNA size fractionation columns (Invitrogen-Life Technologies). The cDNAs derived from each developmental stage of a particular tissue were mixed on an equimolar basis and ligated directionally into the NotI and SalI sites of the pSPOR1 vector (Invitrogen). After ligation of the inserts, the plasmids will be electroporated into DH10B bacteria. Preliminary Library Characterization: Randomly chosen clones from each library were analyzed by restriction digestion to determine average insert size (96 clones) and by sequencing (-4 96-well plates) to confirm library quality (e.g. the presence of short polyA+ tails, genomic DNA contamination (must be <1%), ribosomal RNA clones (must be <1%), etc.] and to provide a sequence database representing the predominant clones in each library. The clones were sequenced at the University of Missouri-Columbia DNA Core Facility. Bioinformatics work was performed by GK Springer's bioinformatics group (WG Spollen, JE Ries, A Guillen, AA Khambati, RV Patel, CM Topinka, SB Bhuiyan) in Computer Science and Health Management and Informatics Departments at the University of Missouri-Columbia. Clone Requests: Requests for clones should be made to the Director of the University of Missouri DNA Core facility at: porcine@rnet.missouri.edu. Citations: 1. Bonaldo MF, Lennon G, Soares MB. Normalization and Subtraction: Two approaches to facilitate gene discovery. Genome Res, 1996; 6:791-806. 2. Jiang H, Bivens NJ, Ries JE, Whitworth KM, Green JA, Forrester LJ, Springer GK, Didion BA, Mathialagan N, Prather RS, Lucy MC (2001) Constructing cDNA libraries with fewer clones that contain long poly(dA) tails. Biotechniques 31:38-42. 3. Soares MB, MF Bonaldo, P Jelene, L Su, L Lawton, A Efstratiadis. 1994. Construction and characterization of a normalized cDNA library. Proc Natl Acad Sci. 91:9228-9232. TAG TISSUE=Day 12 ovarian follicle TAG_SEQ=Not found

ORIGIN

Query Match 84.8%; Score 17.8; DB 7; Length 425;
 Best Local Similarity 90.5%; Pred. No. 2.2e+02;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
 |||||
 Db 49 CCGGGCTGGCGATATGCTAAA 69

RESULT 2

CN153647/c
 LOCUS CN153647 696 bp mRNA linear EST 02-APR-2004
 DEFINITION 940784 MARC 4P1G Sus scrofa cDNA 3', mRNA sequence.
 ACCESSION CN153647
 VERSION CN153647.1 GI:46168077
 KEYWORDS EST.
 SOURCE Sus scrofa (pig)
 ORGANISM Sus scrofa
 Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
 1 (bases 1 to 696)
 Smith,T.P.L., Freking,B.A., Ford,J.J., Vallet,J.L., Wise,T.A.,
 Nonneman,D.J., Wray,J.E. and Keele,J.W.
 Porcine EST collection using a normalized library constructed from embryos representing early developmental stages

```

JOURNAL      Unpublished (2003)
COMMENT      Contact: Smith TPL
             USDA, ARS, US Meat Animal Research Center
             PO Box 166, Clay Center, NE 68933-0166, USA
             Tel: 402 762 4366
             Fax: 402 762 4390
             Email: smith@mail.marc.usda.gov
             Single pass sequencing. Bases called with phred v0.020425.c and
             trimmed with the aid of the trim_alt option. Vector identified with
             cross_match v0.990329.
             Plate: TW8048 row: D column: 8
             Seq primer: TAGAAGCAGCAGTCGAGG.
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             /tissue_type="pooled"
             /lab_host="DH10B"
             /clone_lib="MARC 4PIG"
             /note="Vector: pCDNA3.1; Site 1: EcoRI; Site 2: NotI;
             Library made with combined RNA from day-10, day-13,
             day-15, day-25, and day-30 whole embryos."
ORIGIN
Query Match      84.8%; Score 17.8; DB 7; Length 696;
Best Local Similarity 90.5%; Pred. No. 2.3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
    |||||
Db 488 CCGGGCTGGCGATATGCTAAA 468
    |||||

RESULT 3
LOCUS      CN155761      696 bp      mRNA      linear      EST 02-APR-2004
DEFINITION      943088 MARC 4PIG Sus scrofa cDNA 5', mRNA sequence.
ACCESSION      CN155761
VERSION        CN155761.1 GI:46170191
KEYWORDS       EST.
SOURCE         Sus scrofa (pig)
ORGANISM       Sus scrofa
REFERENCE      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS        Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
TITLE          Smith,T.P.L., Fekking,B.A., Ford,J.J., Vallet,J.L., Wise,T.A.,
               Noneman,D.J., Wray,J.E. and Keeler,J.W.
               Porcine EST collection using a normalized library constructed from
               embryos representing early developmental stages
JOURNAL        Unpublished (2003)
COMMENT        Contact: Smith TPL
               USDA, ARS, US Meat Animal Research Center
               PO Box 166, Clay Center, NE 68933-0166, USA
               Tel: 402 762 4366
               Fax: 402 762 4390
               Email: smith@mail.marc.usda.gov
               Single pass sequencing. Bases called with phred v0.020425.c and
               trimmed with the aid of the trim_alt option. Vector identified with
               cross_match v0.990329.
               Plate: TW8048 row: D column: 8
               Seq primer: GTAATACGACTCACTATAGG.
FEATURES       Location/Qualifiers
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               /tissue_type="pooled"
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               /clone_lib="MARC 4PIG"
               /note="Vector: pCDNA3.1; Site 1: EcoRI; Site 2: NotI;
               Library made with combined RNA from day-10, day-13,
               day-15, day-25, and day-30 whole embryos."
ORIGIN
Query Match      84.8%; Score 17.8; DB 7; Length 696;
Best Local Similarity 90.5%; Pred. No. 2.3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
    |||||
Db 488 CCGGGCTGGCGATATGCTAAA 468
    |||||

RESULT 4
LOCUS      AG570358      998 bp      DNA      linear      GSS 05-JUN-2004
DEFINITION      Mus musculus molossinus DNA, clone:MSMg01-492K10.TJ, genomic survey
               sequence.
ACCESSION      AG570358
VERSION        AG570358.1 GI:48331078
KEYWORDS       GSS.
SOURCE         Mus musculus molossinus
ORGANISM       Mus musculus molossinus
REFERENCE      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS        Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE          Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
               BAC end Sequences of Library MSMg01
JOURNAL        Unpublished
REFERENCE      Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
TITLE          Direct Submission
JOURNAL        Submitted (17-NOV-2003) Masahira Hattori, The Institute of Physical
               and Chemical Research (RIKEN), Genomic Sciences Center (GSC); Japan
               1-7-22 Suehiro-chou,Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan
               (E-mail:hattori@gsc.riken.jp, URL:http://hgp.gsc.riken.go.jp/,
               Tel:81-45-503-9111, Fax:81-45-503-9170)
               Clones are derived from the mouse BAC library MSMg01. For BAC
               library availability, please contact Kuniya Abe (abe@rtc.riken.jp).
               Tsukuba Institute, Bio Resource Center,
               The Institute of Physical and Chemical Research (RIKEN) 3-1-1
               Koyadai, Tsukuba, 305-0074 Japan
               phone: 81-298-36-9189, fax: 81-298-36-9199
               e-mail: abe@rtc.riken.jp
PRIMERS
Sequencing : TJ
LIBRARY      Vector : pBACe3.6
R.Site 1 : EcoRI.
R.Site 2 : EcoRI.
FEATURES     Location/Qualifiers
             1..998
             /organism="Mus musculus molossinus"
             /mol_type="genomic DNA"
             /sub_species="molossinus"
             /db_xref="taxon:57486"
             /clone="MSMg01-492K10.TJ"
             /sex="male"
             /tissue_type="mixture of kidney and spleen"
             /clone_lib="MSMg01 Mouse Male BAC Library"
ORIGIN
Query Match      84.8%; Score 17.8; DB 9; Length 998;
Best Local Similarity 90.5%; Pred. No. 2.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
    |||||
Db 193 CCGGGCTGTCATATCTCTAAA 213
    |||||

RESULT 5
LOCUS      CK699541      429 bp      mRNA      linear      EST 30-MAR-2004
DEFINITION      ZF101-P00082-DBPE-F_G24 GISZF001_ra Danio rerio cDNA clone

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ORIGIN
Query Match      84.8%; Score 17.8; DB 7; Length 696;
Best Local Similarity 90.5%; Pred. No. 2.3e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
    |||||
Db 209 CCGGGCTGGCGATATGCTAAA 229
    |||||

RESULT 4
LOCUS      AG570358      998 bp      DNA      linear      GSS 05-JUN-2004
DEFINITION      Mus musculus molossinus DNA, clone:MSMg01-492K10.TJ, genomic survey
               sequence.
ACCESSION      AG570358
VERSION        AG570358.1 GI:48331078
KEYWORDS       GSS.
SOURCE         Mus musculus molossinus
ORGANISM       Mus musculus molossinus
REFERENCE      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS        Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE          Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
               BAC end Sequences of Library MSMg01
JOURNAL        Unpublished
REFERENCE      Hattori,M., Toyoda,A., Noguchi,H., Kojima,T. and Sakaki,Y.
TITLE          Direct Submission
JOURNAL        Submitted (17-NOV-2003) Masahira Hattori, The Institute of Physical
               and Chemical Research (RIKEN), Genomic Sciences Center (GSC); Japan
               1-7-22 Suehiro-chou,Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan
               (E-mail:hattori@gsc.riken.jp, URL:http://hgp.gsc.riken.go.jp/,
               Tel:81-45-503-9111, Fax:81-45-503-9170)
               Clones are derived from the mouse BAC library MSMg01. For BAC
               library availability, please contact Kuniya Abe (abe@rtc.riken.jp).
               Tsukuba Institute, Bio Resource Center,
               The Institute of Physical and Chemical Research (RIKEN) 3-1-1
               Koyadai, Tsukuba, 305-0074 Japan
               phone: 81-298-36-9189, fax: 81-298-36-9199
               e-mail: abe@rtc.riken.jp
PRIMERS
Sequencing : TJ
LIBRARY      Vector : pBACe3.6
R.Site 1 : EcoRI.
R.Site 2 : EcoRI.
FEATURES     Location/Qualifiers
             1..998
             /organism="Mus musculus molossinus"
             /mol_type="genomic DNA"
             /sub_species="molossinus"
             /db_xref="taxon:57486"
             /clone="MSMg01-492K10.TJ"
             /sex="male"
             /tissue_type="mixture of kidney and spleen"
             /clone_lib="MSMg01 Mouse Male BAC Library"
ORIGIN
Query Match      84.8%; Score 17.8; DB 9; Length 998;
Best Local Similarity 90.5%; Pred. No. 2.4e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAAA 21
    |||||
Db 193 CCGGGCTGTCATATCTCTAAA 213
    |||||

RESULT 5
LOCUS      CK699541      429 bp      mRNA      linear      EST 30-MAR-2004
DEFINITION      ZF101-P00082-DBPE-F_G24 GISZF001_ra Danio rerio cDNA clone

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IMAGE:7165298 5', mRNA sequence.
ACCESSION CK699541
VERSION CK699541.1 GI:42451877
KEYWORDS EST.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
          Cypriniformes; Cyprinidae; Danio.
REFERENCE 1 (bases 1 to 429)
AUTHORS Wei,C., Mathavan,S., Thoreau,H., Lim,L., Lee,C. and Ruan,Y.
TITLE Genome Institute of Singapore, Zebrafish Gene Collection
JOURNAL Unpublished (2004)
COMMENT Contact: Ruan Y
          Cloning and Sequencing
          Genome Institute of Singapore
          60 Biopolis Street, #02-01, Genome, Singapore 138672
          Tel: +65 6478 8073
          Fax: +65 6478 9059
          Email: ruanyj@gis.a-star.edu.sg
          GIS Clone ID: ZF101-P00082-BR2_G24
          PCR Primers
          FORWARD: M13
          BACKWARD: M13
          Plate: ZF101-P00082-BR2 row: G column: 24
          Seq primer: CGCATAACTGTATAGCA
          High quality sequence stop: 429.
FEATURES             source
          Location/Qualifiers
            1..429
               /organism="Danio rerio"
               /mol_type="mRNA"
               /strain="Singapore local strain"
               /db_xref="taxon:7955"
               /clone="IMAGE:7165298"
               /tissue_type="Embryo"
               /dev_stages="7 Different embryonic Stages(From just
               fertilized Embryos to 72 hours just hatched baby fish)"
               /lab_host="DH10B"
               /clone_lib="G15ZF001 ra"
               /note="Vector: pDNR-LIB; Site1: Sfi A (GGCATTACGGCC);
               Site 2: Sfi B (GGCCGAGCGGCC); Priming method: Sfi-(dT)30
               Primed; Priming sequence:
               5.ATTCTAGAGCGCGAGCGCGGACATG(T)30VN ; Directionally
               cloned, 5' cloning site: Sfi A site GGCATTACGGCC ; 5'
               linker/adaptor sequence: 5.AAGCAGTGGTATCAACGAGATGGCC ;
               3' cloning site: Sfi B site GGCGAGCGGCC ; 3'
               linker/adaptor sequence: same as the priming sequence ;
               Average insert size: 2kb ; For PCR insert analysis: Use
               M13 Forward and reverse primers ; Library Amplified ;
               Recombinants (inserts): 98% ; Library complexity: 5x106 ;
               Full-length construction (method): SMART, a Clontech
               method The pooled tissue RNA was collected and used to
               construct full length enriched cDNA library and also
               served as template to synthesize complex first strand cDNA
               probe. Two high density colony arrays were made from over
               110K cDNA clones and hybridized with the probes. Low
               intensity clones were selected as they represented rare
               expressed clones. The hybridization intensities for all
               clones span from 0 to 1.8 million counts and the low
               abundant class ranged from 0 to 13,000."
ORIGIN
Query Match      82.9%; Score 17.4; DB 7; Length 429;
Best Local Similarity 94.7%; Pred. No. 3.6e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 GGGCTGTCGAATATGCTAAA 21
    |||||
Db 3 GGGCTGTCGAATATGCTAAA 21

RESULT 7
CNS000HB 1009 bp DNA linear GSS 03-JUN-1999
Drosophila melanogaster genome survey sequence T7 end of BAC:
BACR35D08 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
AL073822
AL073822.1 GI:4953796
GSS.
KEYWORDS
SOURCE
ORGANISM Drosophila melanogaster (fruit fly)
          Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
          Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
          Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 1009)
AUTHORS Direct Submission
TITLE Genoscope.
JOURNAL Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : sequefgenosco.cns.fr
- Web : www.genoscope.cns.fr)
COMMENT Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see http://www.fruitfly.org The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osogawa and
Aaron Mammoseer in Pieter de Jong's laboratory in the Department of

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LOCUS CB434484
DEFINITION tigr-gss-dog-17000363221477 Dog Library Canis familiaris genomic,
genomic survey sequence.
ACCESSION CB434484
VERSION CB434484.1 GI:36711024
KEYWORDS GSS.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
REFERENCE 1 (bases 1 to 693)
AUTHORS Kirkness,E.F., Bafna,V., Halpern,A.L., Levy,S., Remington,K.,
Rusch,D.B., Delcher,A.L., Pop,M., Wang,W., Fraser,C.M. and
Venter,J.C.
TITLE The dog genome: survey sequencing and comparative analysis
JOURNAL Science 301 (5641), 1898-1903 (2003)
MEDLINE 22875432
PUBMED 14512627
COMMENT Contact: Kirkness EF
          The Institute for Genomic Research
          Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
          Rockville, MD 20850, USA
          Tel: 301-838-0200
          Fax: 301-838-0208
          Email: ekirknes@tigr.org
          Class: shotgun.
          Location/Qualifiers
            1..693
               /organism="Canis familiaris"
               /mol_type="genomic DNA"
               /strain="Standard Poodle"
               /db_xref="taxon:9615"
               /clone_lib="Dog Library"
               /note="Site 1: BstXI; Libraries were prepared from
               peripheral blood"
ORIGIN
Query Match      82.9%; Score 17.4; DB 9; Length 693;
Best Local Similarity 94.7%; Pred. No. 3.8e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 GGGCTGTCGAATATGCTAAA 21
    |||||
Db 326 GGGCTGTCGAATATGCTAAA 308

RESULT 7
CNS000HB 1009 bp DNA linear GSS 03-JUN-1999
Drosophila melanogaster genome survey sequence T7 end of BAC:
BACR35D08 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
AL073822
AL073822.1 GI:4953796
GSS.
KEYWORDS
SOURCE
ORGANISM Drosophila melanogaster (fruit fly)
          Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
          Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
          Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 1009)
AUTHORS Direct Submission
TITLE Genoscope.
JOURNAL Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : sequefgenosco.cns.fr
- Web : www.genoscope.cns.fr)
COMMENT Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see http://www.fruitfly.org The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osogawa and
Aaron Mammoseer in Pieter de Jong's laboratory in the Department of

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Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain Y2; cn bw ep, the same strain used for the BDGP's p1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

1..1009
Location/Qualifiers
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"
/clone="BACR35D08"
/clone_lib="RPCI-98"
/note="end : 7"

ORIGIN

Query Match 81.0%; Score 17; DB 9; Length 1009;
Best Local Similarity 85.7%; Pred. No. 6.4e+02;
Matches 18; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGCTGCTCAATATGCTAAA 21
|||||
Db 133 CCGGCTGCTCAATATGCTAAA 153

RESULT 8

CC888327/c
LOCUS CC888327.1 107 bp DNA linear GSS 31-JUL-2003
DEFINITION SALK_151698.23.35.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_151698.23.35.x, genomic
survey sequence.

ACCESSION CC888327
VERSION CC888327.1 GI:33364847

KEYWORDS GSS.

ORGANISM Arabidopsis thaliana (chale cress)

Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

1 (bases 1 to 107)

Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadrinab,C., Jeake,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished (2001)

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGnAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@salk.edu

This is single pass sequence recovered from the left border of
TDNA.

Class: TDNA tagged.

Location/Qualifiers

1..107

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/ecotype="Col-0"

/db_xref="taxon:3702"

/clone="SALK_151698.23.35.x"

/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN

Query Match 80.0%; Score 16.8; DB 9; Length 107;
Best Local Similarity 90.0%; Pred. No. 6.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CCGGCTGCTCAATATGCTAA 20
|||||
Db 79 CCGGCTGCTCAATATGCTAA 60

RESULT 9

CK091023
LOCUS CK091023 354 bp mRNA linear EST 01-DEC-2003
DEFINITION F039P30.3pR Populus flower cDNA library Populus balsamifera subsp.
trichocarpa cDNA clone F039P30.3, mRNA sequence.

ACCESSION CK091023

VERSION CK091023.1 GI:38575348

KEYWORDS EST.

SOURCE Populus balsamifera subsp. trichocarpa (Populus trichocarpa)

ORGANISM Populus balsamifera subsp. trichocarpa

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

rosids; eurosids I; Malpighiales; Salicaceae; Populus.

1 (bases 1 to 354)

Sterky,F., Bhalarao,R.R., Unneberg,P., Segerman,B., Nilsson,P.,

Brunner,A.M., Campaa,L., Jonsson-Lindvall,J., Tandre,K.,

Strauss,S.H., Sundberg,B., Gustafsson,P., Uhlen,M., Bhalarao,R.P.,

Nilsson,O., Sandberg,G., Karlsson,J., Lundberg,J. and Jansson,S.

A Populus EST resource for functional genomics

Unpublished (2003)

Other_ESTs: F039P30Y, F039P30.5pR

Contact: Bo Segerman

Umea Plant Science Center, Department of Plant Physiology

Umea University

901 87 Umea, Sweden

Tel: +46 90 786 5279

Fax: +46 90 786 6676

Email: bo.segerman@plantphys.umu.se.

Location/Qualifiers

1..354

/organism="Populus balsamifera subsp. trichocarpa"

/mol_type="mRNA"

/sub_species="trichocarpa"

/db_xref="taxon:3694"

/clones="F039P30"

/tissue_type="floral buds"

/clone_lib="Populus flower cDNA library"

/note="Organ: flower"

ORIGIN

Query Match 80.0%; Score 16.8; DB 7; Length 354;
Best Local Similarity 90.0%; Pred. No. 7.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 CCGGCTGCTCAATATGCTAAA 21
|||||
Db 213 CCGGCTTTCAAAATGCTAAA 232

RESULT 10

CK101326/c
LOCUS CK101326 367 bp mRNA linear EST 01-DEC-2003
DEFINITION F039P30.5pR Populus flower cDNA library Populus balsamifera subsp.
trichocarpa cDNA clone F039P30.5, mRNA sequence.

ACCESSION CK101326

VERSION CK101326.1 GI:38585651

KEYWORDS EST.

SOURCE Populus balsamifera subsp. trichocarpa (Populus trichocarpa)

ORGANISM Populus balsamifera subsp. trichocarpa

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

rosids; eurosids I; Malpighiales; Salicaceae; Populus.

1 (bases 1 to 367)

AUTHORS Sterky, F., Bhalarao, R.R., Unneberg, P., Segerman, B., Nilsson, P., Brunner, A.M., Campaa, L., Jonsson-Lindvall, J., Tandare, K., Straus, S.H., Sundberg, B., Gustafsson, P., Uhlen, M., Bhalarao, R.P., Nilsson, O., Sandberg, G., Karlsson, J., Lundberg, J., and Jansson, S. A Populus EST resource for functional genomics

TITLE Unpublished (2003)

JOURNAL

COMMENT Other ESTs: F039P30V, F039P30.3P

Contact: Bo Segerman

Umea Plant Science Center, Department of Plant Physiology

Umea University

901 87 Umea, Sweden

Tel: +46 90 786 5279

Fax: +46 90 786 6676

Email: bo.segerman@plantphys.umu.se.

Location/Qualifiers

1..367

/organism="Populus balsamifera subsp. trichocarpa"

/mol_type="mRNA"

/sub_species="trichocarpa"

/db_xref="taxon:3694"

/clone="F039P30"

/tissue_type="floral buds"

/clone_lib="Populus flower cDNA library"

/note="Organ: flower"

ORIGIN

Query Match 80.0%; Score 16.8; DB 7; Length 367;

Best Local Similarity 90.0%; Pred. No. 7.4e+02;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 CCGGCTGTCATATGCTAA 21

||||| ||||| ||||| ||||| |||||

Db 318 CCGGCTTCAGATGCTAA 299

||||| ||||| ||||| ||||| |||||

RESULT 11

BG037611

LOCUS BG037611.1 GI:12480196

DEFINITION BG037611.1 407 bp mRNA linear EST 24-JAN-2001

VERSION BG037611.1

KEYWORDS EST.

SOURCE Xenopus laevis (African clawed frog)

ORGANISM Xenopus laevis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae; Xenopodinae; Xenopus; Xenopus.

1 (bases 1 to 407)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

Unpublished (1997)

Other ESTs: dc53e09.x1

Contact: Robert Strausberg, Ph.D.

Email: cgapbs@mail.nih.gov

Tissue Procurement: Martha Rebert, Steven L. Klein, Ph.D.

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

Cloning Sequencing by: Washington University Genome Sequencing Center

Clone distribution: Xenopus clones from this library are available through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov

Seq primer: -40RP from Gibco

High quality sequence stop: 403.

Location/Qualifiers

1..407

/organism="Xenopus laevis"

/mol_type="mRNA"

/db_xref="taxon:8355"

/clones="IMAGE:3400816"

/tissue_type="embryo (stages 24-25)"

/lab_host="DH10B (phage-resistant)"

/clone_lib="NICHD_XGC_Emb3"

FEATURES

source

1..501

/organism="Malus x domestica"

/mol_type="mRNA"

/db_xref="taxon:3750"

/clones="Mdfw2019108"

/lab_host="DH10B ampicillin resistant"

/clone_lib="Mdfw"

/note="Vector: DH10B ampicillin resistant; Site_1: NotI; Site_2: EcoRII; Total RNA was extracted separately from each stage (bud, balloon, open and after pollination), using the 'pine tree' method. Poly(A)+mRNA was isolated twice from total RNA from each stage using the Oligotex Direct mRNA kit (Qiagen). mRNA was reverse transcribed into double stranded cDNA using a modified oligo(dT) primer with an identifying tag sequence (see table below). cDNAs from different stages were pooled in equal amounts before adaptor ligation. Tag identification when sequencing from 5' end: Stage 1 (bud) insert 18(A)TCGGA; Stage 2 (balloon) insert 18(A)TCGGA; Stage 3 (open) insert 18(A)TCGCT; Stage 4 (after pollination) insert 18(A)TCGGT. Tag identification when sequencing from 3' end: Stage 1 (bud) TCCGAl8(T) insert; Stage 2 (balloon) TCGGAl8(T) insert; Stage 3 (open) ACGGAl8(T) insert; Stage 4 (after

/note="Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.7 kb. Constructed by Life Technologies. Note: This is a Xenopus Gene Collection (XGC) library."

ORIGIN

Query Match 80.0%; Score 16.8; DB 4; Length 407;

Best Local Similarity 85.7%; Pred. No. 7.4e+02;

Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 CCGGGCTGTCATATGCTAA 21

||||| ||||| ||||| ||||| |||||

Db 364 CAGGGTGTNAATATGCTAA 384

||||| ||||| ||||| ||||| |||||

RESULT 12

CM489600/c

LOCUS CM489600.1 GI:46603708

DEFINITION CM489600.1 501 bp mRNA linear EST 24-MAY-2004

VERSION CM489600.1

KEYWORDS EST.

SOURCE Malus x domestica (cultivated apple)

ORGANISM Malus x domestica

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids 1; Rosales; Rosaceae; Maloideae; Malus.

1 (bases 1 to 501)

Korban, S., Vodkin, L., Liu, L., Gasic, K., Gonzales, O., Hernandez, A., Aldwinckle, H., Malnoy, M., Carroll, N., Goldebrogh, P., Orvis, K., Clifton, S., Pape, D., Marra, M., Hillier, L., Martin, J., Wylie, T., Dante, M., Theising, B., Bowers, Y., Gibbons, M., Ritter, E., Ronko, I., Tsagaris, R., Kennedy, S., Waterston, R., and Wilson, R.

Apple Functional Genomics grant - NSF 0321702

Unpublished (2004)

Contact: Schuyler S. Korban

Apple Functional Genomics grant - NSF 0321702

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

Library materials provided by: Schuyler S. Korban Library

constructed by: A. Hernandez / K. Gasic Library sequenced by: Washington University Genome Sequencing Center

WASHU EST name: aaf76f04.y1

High quality sequence stop: 425.

Location/Qualifiers

1..501

/organism="Malus x domestica"

/mol_type="mRNA"

/db_xref="taxon:3750"

/clones="Mdfw2019108"

/lab_host="DH10B ampicillin resistant"

/clone_lib="Mdfw"

/note="Vector: DH10B ampicillin resistant; Site_1: NotI; Site_2: EcoRII; Total RNA was extracted separately from each stage (bud, balloon, open and after pollination), using the 'pine tree' method. Poly(A)+mRNA was isolated twice from total RNA from each stage using the Oligotex Direct mRNA kit (Qiagen). mRNA was reverse transcribed into double stranded cDNA using a modified oligo(dT) primer with an identifying tag sequence (see table below). cDNAs from different stages were pooled in equal amounts before adaptor ligation. Tag identification when sequencing from 5' end: Stage 1 (bud) insert 18(A)TCGGA; Stage 2 (balloon) insert 18(A)TCGGA; Stage 3 (open) insert 18(A)TCGCT; Stage 4 (after pollination) insert 18(A)TCGGT. Tag identification when sequencing from 3' end: Stage 1 (bud) TCCGAl8(T) insert; Stage 2 (balloon) TCGGAl8(T) insert; Stage 3 (open) ACGGAl8(T) insert; Stage 4 (after

pollination) ACCGA18(T) insert. Double stranded cDNAs were size selected (more than 450 bp), adapted with EcoRI adaptors at both ends and then digested with NotI. The cDNAs were then directionally cloned into EcoRI-NotI digested pBS II SK(+) phagemid vector(Stratagene). Identification of adaptors and tags in 5'-end sequenced clones: <vector>...TAGCTTC<End Vector><Start EcoRI adaptor>GATATCGAATTCCTATTGTGGC<End EcoRI adaptor>Start Insert<...AAAAAAAAAAAAAAAA<End Insert><Start Tag>TGCAG<End Tag><Start NotI site/Vector>GGCGCGCCACGCGG... The total number of white colony forming units (cfu) in the primary library before amplification was 1.1x10⁶ cfu (colony forming units). The background of empty clones was less than 1%. Inserts ranged from 0.5 kb to 3 kb, as determined by PCR. Purified plasmid DNA from the primary library was converted to single-stranded circles and used as a template for PCR amplification using the T7 and T3 priming sites flanking the cloned cDNA inserts. The purified PCR products, representing the entire cloned cDNA population, were used as a driver for normalization. Hybridization between the single-stranded library and the PCR products was carried out for 4 hours at 30C. Unhybridized single-stranded DNA circles were separated from hybridized DNA rendered partially double-stranded and electroporated into DH10B cells to generate the normalized library. The total number of clones with insert was 9x10⁶ cfu. Background of empty clones was less than 1%.

ORIGIN

| | | | | |
|-----------------------|-------|--------------------|-------|-------------|
| Query Match | 80.0% | Score 16.8; | DB 7; | Length 501; |
| Best Local Similarity | 85.7% | Pred. No. 7.6e+02; | | |

QY 1 CCGGGCTGTCAATATGCTAAA 21
|||
332 CCGGGCTGTCCATTTGCNAAA 312
Db

RESULT 13

| AZ981379/c | | 516 bp | DNA | linear | GSS 27-APR-2001 |
|------------|---|--------|-----|--------|-----------------|
| LOCUS | AZ981379 | | | | |
| DEFINITION | 2M0258113R Mouse 10kb plasmid UUGC2M library Mus musculus genomic clone UUGC2M0258113 R, genomic survey sequence. | | | | |

FEATURES

source

1. .516
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0258113"
/sex="Female"
/lab_host="E. coli strain XL10-Gold, Tl-resistant, F-"
/clone_lib="Mouse 10kb plasmid UUGC2M library"
/note="vector: PWD42nb; Purified genomic DNA from M.
musculus C57BL/6J (female) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptored DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 [gil473214|gb|AF129072.1], a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptored mouse DNA was annealed to
adaptored vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

ORIGIN

| | | | | |
|-----------------------|-----------------|--------------------|-----------|-------------|
| Query Match | 80.0%; | Score 16.8; | DB 8; | Length 516; |
| Best Local Similarity | 90.0%; | Pred. No. 7.6e+02; | | |
| Matches 18; | Conservative 0; | Mismatches 2; | Indels 0; | Gaps 0; |

QY 2 CGGGCTGTCAATATGCTAAA 21
Dp 30 CTGGCTGTCAATATGCTACA 11

RESULT 14

| BU815985/c | BU815985 | 562 bp | mRNA | linear | EST 15-OCT-2002 |
|------------|----------|--------------|--------------|---------------------------------------|------------------------------|
| LOCUS | N058F04 | Populus bark | CDNA library | Populus tremula x Populus tremuloides | cdna 5 prime, mRNA sequence. |
| DEFINITION | | | | | |

FEATURES

```

source
1. 562
/organism="Populus tremula x Populus tremuloides"
/mol_type="mRNA"
/db_xref="taxon:47664"
/tissue_type="bark"
/clone_lib="Populus bark cDNA library"

```

ORIGIN

Query Match 80.0%; Score 16.8; DB 5; Length 562;
 Best Local Similarity 90.0%; Pred. No. 7.7e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 CGGGCTGTCAATATGCTAAA 21
 ||||| ||||| ||||| ||||| |||||
 Db 293 CGGGCTTTCAGATGCTAAA 274

RESULT 15
 CAB21839/c
 LOCUS
 DEFINITION RSH08C06 two-month-old roots from clone 'Beaupre' grown for 19 days under restricted irrigation Populus balsamifera subsp. trichocarpa x Populus deltoides cDNA 5', mRNA sequence.

ACCESSION CAB21839
 VERSION CAB21839.1 GI:28605388
 KEYWORDS EST.
 SOURCE Populus balsamifera subsp. trichocarpa x Populus deltoides
 ORGANISM Populus balsamifera subsp. trichocarpa x Populus deltoides
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids I; Malpighiales; Salicaceae; Populus.

REFERENCE 1 (bases 1 to 620)
 Kohler,A., Delaruelle,C., Martin,D., Encelot,N. and Martin,P.
 The poplar root transcriptome: analysis of 7000 expressed sequence tags
 FEBS Lett. 542 (1-3), 37-41 (2003)
 CONTACT: Martin FM
 Institut de Microbiologie Forestiere
 Institut National de la Recherche Agronomique
 Centre INRA de Nancy, 54280 Champenoux, France
 Tel: +33 383 39 40 80
 Fax: +33 383 39 40 89
 Email: fmartin@nancy.inra.fr
 Insert Length: 620 Std Error: 0.00
 Seq primer: Forwat 5' AAGCGCGCATTGTGTGGTACCC.

FEATURES
 source
 1..620
 /organism="Populus balsamifera subsp. trichocarpa x Populus deltoides"
 /mol_type="mRNA"
 /cultivar="Beaupre"
 /db_xref="taxon:3695"
 /dev_stage="two-month-old"
 /clone_lib="two-month-old roots from clone 'Beaupre' grown for 19 days under restricted irrigation"
 /note="Organ: root; Vector: pTriplEx2; cDNA library of roots from two-month-old Populus trichocarpa Torr. & Gray x deltoides Bartr. Ex Marshall (clone 'Beaupre') grown for 19 days under restricted irrigation to reach 50% of the transpiration rate of fully watered plants. The cDNA library was constructed from 1 ug of total RNA using the SMART cDNA synthesis kit (Clontech, Palo Alto, CA, USA) according to the manufacturer's instructions. The resulting cDNA was packed into lambda phages using the Gigapack III Gold packaging kit (Stratagene, La Jolla, CA). The pTriplEx2 phagemid clones in Escherichia coli were obtained by using the mass in vivo excision protocol according to the manufacturer's instructions (Clontech)."

ORIGIN
 Query Match 80.0%; Score 16.8; DB 6; Length 620;
 Best Local Similarity 90.0%; Pred. No. 7.8e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 CGGGCTGTCAATATGCTAAA 21
 ||||| ||||| ||||| ||||| |||||
 Db 389 CGGGCTTTCAGATGCTAAA 370

RESULT 16
 A2832056

LOCUS
 DEFINITION A2832056 623 bp DNA linear GSS 20-FEB-2001
 Clone UUSC2M0112L09 F, genomic survey sequence.

ACCESSION A2832056
 VERSION A2832056.1 GI:13001964
 KEYWORDS GSS.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 623)
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Ielam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.,
 Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
 Unpublished (2000)
 CONTACT: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0112 row: L column: 09
 Seq primer: CGTGTAAACGACGCCAGT
 Class: plasmid ends
 High quality sequence stop: 623.

FEATURES
 source
 1..623
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUSC2M0112L09"
 /sex="Male"
 /lab_hosts="E. Coli strain XL10-Gold, Tl-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUSC1M library"
 /note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 (gi|4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN
 Query Match 80.0%; Score 16.8; DB 8; Length 623;
 Best Local Similarity 90.0%; Pred. No. 7.8e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 CGGGCTGTCAATATGCTAAA 21
 ||||| ||||| ||||| ||||| |||||
 Db 563 CTGGCTGTCAATATGCTACA 582

RESULT 17
 CK317796/c


```

LOCUS       CK317796               631 bp    mRNA    linear    EST 11-MAY-2004
DEFINITION   B9P01h01 Populus stem seasonal library Populus deltoides cDNA, mRNA
sequence.
ACCESSION   CK317796
VERSION     CK317796.1   GI:47106219
KEYWORDS    EST.
SOURCE      Populus deltoides
ORGANISM    Populus deltoides
REFERENCE   1 (bases 1 to 631)
AUTHORS     Park, S. and Han, K.-H.
TITLE       Gene expression profile during seasonal growth cycle in poplar tree
JOURNAL     Unpublished (2003)
COMMENT     Contact: Kyung-Hwan Han
            Department of Forestry
            Michigan State University
            126 Natural Resources, East Lansing, MI 48824-1222, USA
            Tel: 517 353 4751
            Fax: 517 432 1143
            Email: hanky@msu.edu.

FEATURES             Location/Qualifiers
     source          1..631
                     /organism="Populus deltoides"
                     /mol_type="mRNA"
                     /strain="ILL-129"
                     /db_xref="taxon:3696"
                     /tissue_type="stem"
                     /dev_stages="1 year old"
                     /clone_lib="Populus stem seasonal library"

ORIGIN
Query Match      80.0%; Score 16.8; DB 7; Length 631;
Best Local Similarity 90.0%; Pred. No. 7.8e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Cy  2  CGGGCTGTCAATATGCTAAA 21
      ||||| ||||| ||||| |||||
Db   355 CGGGCTTTCAAGATGCTAAA 336

RESULT 18
BU863468/c
LOCUS       BU863468               635 bp    mRNA    linear    EST 16-OCT-2002
DEFINITION   S028D11 Populus imbibed seed cDNA library Populus tremula cDNA 5
prime, mRNA sequence.
ACCESSION   BU863468
VERSION     BU863468.1   GI:24049528
KEYWORDS    EST.
SOURCE      Populus tremula
ORGANISM    Populus tremula
REFERENCE   1 (bases 1 to 635)
AUTHORS     Unneberg, P., Bhalerao, R.R., Jansson, S. and Sterky, F.
TITLE       The poplar tree transcriptome: Analysis of expressed sequence tags
            from multiple libraries
JOURNAL     Unpublished (2002)
COMMENT     Contact: BHALERAO RUPALI R.
            Umea Plant Science Center
            Department of Plant Physiology
            University of Umea, 901 87 Umea, Sweden
            Tel: +46 90 786 5279
            Fax: +46 90 786 6676
            Email: rupali.bhalerao@plantphys.umu.se.

FEATURES             Location/Qualifiers
     source          1..635
                     /organism="Populus tremula"
                     /mol_type="mRNA"
                     /db_xref="taxon:113636"
                     /tissue_type="imbibed seed"

```

```

/clone_lib="Populus imbibed seed cDNA library"

Query Match      80.0%; Score 16.8; DB 5; Length 635;
Best Local Similarity 90.0%; Pred. No. 7.8e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Cy  2  CGGGCTGTCAATATGCTAAA 21
      ||||| ||||| ||||| |||||
Db   340 CGGGCTTTCAAGATGCTAAA 321

RESULT 19
CV257160
LOCUS       CV257160               751 bp    mRNA    linear    EST 22-SEP-2004
DEFINITION   WS0245.B21 N14 PTx-D-ICC-N-A-14 Populus balsamifera subsp.
            trichocarpa x Populus deltoides cDNA clone WS0245_N14 3', mRNA
            sequence.
ACCESSION   CV257160
VERSION     CV257160.1   GI:52510135
KEYWORDS    EST.
SOURCE      Populus balsamifera subsp. trichocarpa x Populus deltoides
ORGANISM    Populus balsamifera subsp. trichocarpa x Populus deltoides
REFERENCE   1 (bases 1 to 751)
AUTHORS     Ralph, S., Cooper, D., Kolosova, N., Oddy, C., Butterfield, Y.,
            Kirkpatrick, R., Liu, J., Palmquist, D., Stott, J., Barber, S., Yang, G.,
            Babakaiff, R., Brown-John, M., Chand, S., Featherstone, R., Mason, A.,
            Mayo, M., Moran, J., Olson, F., Wong, D., Ritland, C.E., Siddiqui, A., and
            Holt, R., Jones, S., Marra, M., Ellis, B.E., Douglas, C., Ritland, K. and
            Bohlmann, J.
TITLE       The poplar transcriptome: Analysis of expressed sequence tags from
            multiple cDNA libraries
JOURNAL     Unpublished (2004)
COMMENT     Contact: Joerg Bohlmann
            Genome BC forest genomics program
            University of British Columbia
            UBC Biotechnology Laboratory, 6174 University Boulevard, Rm. 237,
            Vancouver, British Columbia, Canada, V6T 1Z3
            Tel: 1-604-822-0282
            Fax: 1-604-822-6097
            Email: bohlmann@interchange.ubc.ca
            Plate: WS0245 row: N column: 14
            High quality sequence stop: 751
            POLYA=Yes.

FEATURES             Location/Qualifiers
     source          1..751
                     /organism="Populus balsamifera subsp. trichocarpa x
                     Populus deltoides"
                     /mol_type="mRNA"
                     /cultivar="H11-11"
                     /db_xref="taxon:3695"
                     /clone="WS0245_N14"
                     /sex="Male"
                     /lab_host="E. coli DH10B T1 phage resistant cells"
                     /clone_lib="PTx-D-ICC-N-A-14"
                     /note="Vector: pBluescript II SK (+) XR; Site 1: EcoRI (5'
                     end of cDNA); Site 2: XhoI (3' end of cDNA); Cultured
                     cells (de Sa MM et al. (1992) Plant Physiology 98:728-737)
                     were grown in media (45mL) supplemented with either 50uM
                     salicylic acid, 50uM benzothiadiazole, 50uM methyl
                     jasmonate, 20ug chitosan or 200uL of Pollacia radiosa
                     extract. Cells were harvested after a 3 hour treatment,
                     along with untreated control cells. mRNA was isolated from
                     each tissue source independently and equal quantities of
                     mRNA from each tissue were then pooled. cDNA was prepared
                     from 5 micrograms of mRNA and directionally ligated into
                     the pBluescript II SK (+) XR vector using the pBluescript
                     II XR cDNA Library Construction Kit according to
                     manufacturer's instructions with modifications
                     (Stratagene). Plasmid DNA was then transformed by

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electroporation into DH10B cells (Invitrogen) for propagation. Normalization was applied according to published methods [Bonaldo M.F. et al. (1996) Genome Research 6(9):791] in order to reduce the abundance of highly expressed transcripts."

ORIGIN

Query Match 80.0%; Score 16.8; DB 7; Length 751;
 Best Local Similarity 90.0%; Pred. No. 8e+02; 2; Indels 0; Gaps 0;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 CGGGCTGTCAATATGCTAAA 21
 |||||
 Db 491 CGGGCTTTCAGATGCTAAA 510

RESULT 20

CR268537/c
 LOCUS CR268537 790 bp DNA linear GSS 06-JUL-2004
 DEFINITION Reverse strand read from insert in 5'HPRT insertion targeting and chromosome engineering clone MHPN344a20, genomic survey sequence.
 ACCESSION CR268537.1 GI:50047390
 VERSION GSS; genome survey sequence; MICR.
 KEYWORDS Mus musculus (house mouse)
 SOURCE Mus musculus
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 790)
 REFERENCE Adams, D.J., Biggs, P.J., Cox, A.V., Davies, R.M., van der Weyden, L., Jonkers, J., Smith, J., Plumb, R.W., Taylor, R.G., Nishijima, I., Yu, Y., Rogers, J. and Bradley, A.
 AUTHORS Direct Submission
 TITLE Submitted (20-FEB-2004) Sanger Centre, Hinxton, Cambridgeshire,
 JOURNAL CB10 18A, UK. http://www.sanger.ac.uk/MICR
 FEATURES
 Location/Qualifiers
 1..790
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /db_xref="taxon:10090"
 /clone="MHPN344a20"
 /clone_lib="MHPN"

ORIGIN

Query Match 80.0%; Score 16.8; DB 9; Length 790;
 Best Local Similarity 90.0%; Pred. No. 8e+02; 2; Indels 0; Gaps 0;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 CGGGCTGTCAATATGCTAAA 21
 |||||
 Db 361 CTGGCTGTCAATATGCTACA 342

RESULT 21

CR016265/c
 LOCUS CR016265 900 bp DNA linear GSS 05-JUL-2004
 DEFINITION Forward strand read from insert in 3'HPRT insertion targeting and chromosome engineering clone MHPpd17, genomic survey sequence.
 ACCESSION CR016265
 VERSION GSS; genome survey sequence; MICR.
 KEYWORDS Mus musculus (house mouse)
 SOURCE Mus musculus
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 900)
 REFERENCE Adams, D.J., Biggs, P.J., Cox, A.V., Davies, R.M., van der Weyden, L., Jonkers, J., Smith, J., Plumb, R.W., Taylor, R.G., Nishijima, I., Yu, Y., Rogers, J. and Bradley, A.
 AUTHORS Direct Submission
 TITLE Submitted (20-FEB-2004) Sanger Centre, Hinxton, Cambridgeshire,
 JOURNAL CB10 18A, UK. http://www.sanger.ac.uk/MICR
 FEATURES
 Location/Qualifiers

source

1..900
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /db_xref="taxon:10090"
 /clone="MHPpd17"
 /clone_lib="MHP"

ORIGIN

Query Match 80.0%; Score 16.8; DB 9; Length 900;
 Best Local Similarity 90.0%; Pred. No. 8.1e+02; 2; Indels 0; Gaps 0;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 CCGGGCTGTCAATATGCTAA 20
 |||||
 Db 145 CCGGGCTGTCAAGATGCTCA 164

RESULT 22

BQ220271/c
 LOCUS BQ220271 986 bp mRNA linear EST 02-MAY-2002
 DEFINITION AGENCOURT 7572589 NIH_MGC_92 Homo sapiens cDNA clone IMAGE:6044520 5', mRNA sequence.
 ACCESSION BQ220271
 VERSION BQ220271.1 GI:20401671
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 986)
 REFERENCE NIH-MGC http://mgc.nci.nih.gov/.
 AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE Unpublished (1999)
 JOURNAL
 COMMENT Contact: Robert Straubeberg, Ph.D.
 Email: cgabbs-remail.nih.gov
 Tissue Procurement: ATCC
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
 Plate: LLAM13287 row: e column: 01
 High quality sequence stop: 149.

FEATURES

Location/Qualifiers
 1..986
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:6044520"
 /tissue_type="embryonal carcinoma, cell line"
 /lab_host="DH10B (phage-resistant)"
 /clone_lib="NIH_MGC_92"
 /note="Organ: testis; Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI; Cloned unidirectionally; oligo-dT primed. Average insert size 2.5 kb. Library enriched for full-length clones and constructed by Life Technologies. Note: this is a NIH_MGC Library."

ORIGIN

Query Match 80.0%; Score 16.8; DB 5; Length 986;
 Best Local Similarity 90.0%; Pred. No. 8.2e+02; 2; Indels 0; Gaps 0;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 2 CCGGGCTGTCAATATGCTAAA 21
 |||||
 Db 758 CAGGCTGTCGATATGCTAAA 739

RESULT 23

CE199293/c
 LOCUS CE199293 298 bp DNA linear GSS 25-SEP-2003
 DEFINITION tigr-gss-dog-17000372211318 Dog Library Canis familiaris genomic,

```

genomic survey sequence.
ACCESSION CE199293
VERSION CE199293.1 GI:35354946
KEYWORDS GSS.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
REFERENCE 1 (bases 1 to 298)
AUTHORS Kirkness,E.F., Bafna,V., Halpern,A.L., Levy,S., Remington,K.,
Rusch,D.B., Delcher,A.L., Pop,M., Wang,W., Fraser,C.M. and
Venter,J.C.
The dog genome: survey sequencing and comparative analysis
JOURNAL Science 301 (5641), 1898-1903 (2003)
MEDLINE 22875432
PUBMED 14512627
COMMENT Contact: Kirkness EF
The Institute for Genomic Research
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
Rockville, MD 20850, USA
Tel: 301-838-0200
Fax: 301-838-0208
Email: ekirknes@tigr.org
Class: shotgun.
FEATURES             Location/Qualifiers
     source           1..298
                     /organism="Canis familiaris"
                     /mol_type="genomic DNA"
                     /strain="Standard Poodle"
                     /db_xref="taxon:9615"
                     /clone_lib="Dog Library"
                     /note="Site 1: BatXI; Libraries were prepared from
                     peripheral blood"
ORIGIN
Query Match       78.1%; Score 16.4; DB 9; Length 298;
Best Local Similarity 94.4%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      4  GGCTGTCATATGCTAAA 21
        ||||| ||||| |||||
Db      241 GGCTGTAATATGCTAAA 258

RESULT 24
AL926049/c
LOCUS      AL926049      363 bp      mRNA      linear      EST 06-JUL-2004
DEFINITION AL926049 PJR-Z1-Z2 Danio rerio cDNA clone 164-D08-2, mRNA sequence.
ACCESSION AL926049
VERSION AL926049.1 GI:23192629
KEYWORDS EST.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
REFERENCE 1 (bases 1 to 363)
AUTHORS Lo,J., Lee,S., Xu,M., Liu,F., Ruan,H., Eun,A., He,Y., Ma,W.,
Wang,W., Wen,Z. and Peng,J.
15000 unique zebrafish EST clusters and their future use in
microarray for profiling gene expression patterns during
embryogenesis
JOURNAL Genome Res. 13 (3), 455-466 (2003)
MEDLINE 22505427
PUBMED 12618376
COMMENT Contact: Peng J
Lab of Functional Genomics
Institute of Molecular and Cell Biology
30 Medical Drive, Singapore, 117609, Singapore
Email: pengjr@imcb.a-star.edu.sg
Clone requests: info@openbiosystems.com
Open Biosystems,
6705 Odyssey Drive, Huntsville, AL 35806.

```

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FEATURES             Location/Qualifiers
     source           1..363
                     /organism="Danio rerio"
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                     /db_xref="taxon:7955"
                     /clone_lib="164-D08-2"
                     /tissue_type="whole embryo or fish"
                     /dev_stage="mixed stages"
                     /clone_lib="PJR-Z1-Z2"
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Best Local Similarity 94.4%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      4  GGCTGTCATATGCTAAA 21
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Db      239 GGCTGTCAGATGCTAAA 222

RESULT 25
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LOCUS      AL152957      502 bp      mRNA      linear      EST 19-APR-2001
DEFINITION LB45125.Sprime LD Drosophila melanogaster embryo pOT2 Drosophila
melanogaster cDNA clone LD45125 Sprime similar to D83486: Su(fu)
FBgn0005355 PID:gl1208417 SPTREMBL:Q27279, mRNA sequence.
ACCESSION AL152957
VERSION AL152957.1 GI:4422375
KEYWORDS EST.
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 502)
AUTHORS Harvey,D., Brokstein,P., Hong,L., Evans-Holm,M., Su,C., Tsang,G.,
Lewis,S. and Rubin,G.M.
BDGP/HHMI Drosophila EST Project
Unpublished (2001)
COMMENT Contact: Stapleton, M.
BDGP
Lawrence Berkeley National Lab
One Cyclotron Rd, Berkeley, CA 94720, USA
Fax: 510 486 6798
Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
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ORIGIN
Query Match       78.1%; Score 16.4; DB 1; Length 502;
Best Local Similarity 94.4%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1  CCGGGCTGTCATATGCT 18
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Search completed: September 6, 2005, 21:55:48

Job time : 1507.84 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 20:39:40 ; Search time 1492 Seconds
(without alignments)
746.964 Million cell updates/sec

Title: US-10-729-421-8

Perfect score: 23

Sequence: 1 tcattgactgcaattccgtcttt 23

Scoring table: OLIGO NUC

Gapop_60.0 , Gapext 60.0

Searched: 4708233 seqs, 24227607955 residues

Word size : 10

Total number of hits satisfying chosen parameters: 417

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Listing first 45 summaries

Database :

GenEmbl.*

1: gb_ba.*

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4: gb_cm.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_ro.*

11: gb_sbs.*

12: gb_sy.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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| C 1 | 20 | 87.0 | 36 | AX589722 | AX589722 Sequence |
| C 2 | 20 | 87.0 | 39 | AX589721 | AX589721 Sequence |
| C 3 | 19 | 82.6 | 33 | AX589711 | AX589711 Sequence |
| C 4 | 19 | 82.6 | 36 | AX589702 | AX589702 Sequence |
| C 5 | 19 | 82.6 | 56 | AX224249 | AX224249 Sequence |
| 6 | 12 | 52.2 | 25 | AX182204 | AX182204 Sequence |
| 7 | 12 | 52.2 | 25 | AX382013 | AX382013 Sequence |
| 8 | 12 | 52.2 | 34 | BD173847 | BD173847 JNK inhib |
| 9 | 12 | 52.2 | 51 | AR444320 | AR444320 Sequence |
| 10 | 12 | 52.2 | 51 | AR444321 | AR444321 Sequence |
| 11 | 12 | 52.2 | 60 | CQ536043 | CQ536043 Sequence |
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| C 13 | 11 | 47.8 | 15 | AR119503 | AR119503 Sequence |
| C 14 | 11 | 47.8 | 16 | AR285636 | AR285636 Sequence |
| C 15 | 11 | 47.8 | 16 | AR397627 | AR397627 Sequence |
| 16 | 11 | 47.8 | 20 | AR167035 | AR167035 Sequence |
| 17 | 11 | 47.8 | 20 | AR210690 | AR210690 Sequence |
| C 18 | 11 | 47.8 | 20 | AR301418 | AR301418 Sequence |
| C 19 | 11 | 47.8 | 20 | AR313575 | AR313575 Sequence |

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| 22 | 11 | 47.8 | 21 | 6 | AR210652 | AR210652 Sequence |
| 23 | 11 | 47.8 | 22 | 6 | CQ768639 | CQ768639 Sequence |
| 24 | 11 | 47.8 | 24 | 6 | AX935032 | AX935032 Sequence |
| C 25 | 11 | 47.8 | 25 | 6 | AR364418 | AR364418 Sequence |
| C 26 | 11 | 47.8 | 25 | 6 | AR568240 | AR568240 Sequence |
| C 27 | 11 | 47.8 | 26 | 6 | AR542624 | AR542624 Sequence |
| C 28 | 11 | 47.8 | 26 | 6 | AX235895 | AX235895 Sequence |
| C 29 | 11 | 47.8 | 26 | 6 | AX402749 | AX402749 Sequence |
| C 30 | 11 | 47.8 | 27 | 6 | BD183051 | BD183051 Nucleic a |
| C 31 | 11 | 47.8 | 27 | 6 | I22149 | I22149 Sequence 8 |
| C 32 | 11 | 47.8 | 28 | 6 | I13959 | I13959 Sequence 38 |
| 33 | 11 | 47.8 | 30 | 6 | AX085450 | AX085450 Sequence |
| 34 | 11 | 47.8 | 30 | 6 | AX101005 | AX101005 Sequence |
| 35 | 11 | 47.8 | 33 | 6 | AR004393 | AR004393 Sequence |
| C 36 | 11 | 47.8 | 33 | 6 | AR005205 | AR005205 Sequence |
| 37 | 11 | 47.8 | 33 | 6 | AR005206 | AR005206 Sequence |
| C 38 | 11 | 47.8 | 33 | 6 | AR064955 | AR064955 Sequence |
| C 39 | 11 | 47.8 | 33 | 6 | AR072936 | AR072936 Sequence |
| C 40 | 11 | 47.8 | 33 | 6 | AR072938 | AR072938 Sequence |
| 41 | 11 | 47.8 | 33 | 6 | AR097185 | AR097185 Sequence |
| 42 | 11 | 47.8 | 33 | 6 | AR130683 | AR130683 Sequence |
| 43 | 11 | 47.8 | 33 | 6 | AR172032 | AR172032 Sequence |
| 44 | 11 | 47.8 | 33 | 6 | BD189149 | BD189149 HCV Genom |
| 45 | 11 | 47.8 | 33 | 6 | BD189296 | BD189296 HCV Genom |

ALIGNMENTS

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| RESULT 1 | AX589722/c | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| LOCUS | AX589722/c | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| DEFINITION | AX589722/c | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| ACCESSION | AX589722 | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| VERSION | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| KEYWORDS | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| SOURCE | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| ORGANISM | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| REFERENCE | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| AUTHORS | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| TITLE | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| JOURNAL | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| FEATURES | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| source | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
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| /mol_type="unassigned DNA" | AX589722.1 | GI:27901012 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
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| Query Match | AX589722 | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| Best Local Similarity | AX589722 | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| Matches | AX589722 | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
| Conservative | AX589722 | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
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| 0; | AX589722 | AX589722 | Sequence 29 from Patent WO02081621. | 36 bp | DNA | linear | PAT 24-JAN-2003 |
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| Db | 31 | TCATGACTGCAATTCGGTC | 12 | TCATGACTGCAATTCGGTC | 12 | TCATGACTGCAATTCGGTC | 12 |
| RESULT 2 | AX589721/c | AX589721 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| LOCUS | AX589721/c | AX589721 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| DEFINITION | AX589721/c | AX589721 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| ACCESSION | AX589721 | AX589721 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| VERSION | AX589721.1 | GI:27901011 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| KEYWORDS | AX589721.1 | GI:27901011 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| SOURCE | AX589721.1 | GI:27901011 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| ORGANISM | AX589721.1 | GI:27901011 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| synthetic construct | AX589721.1 | GI:27901011 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| synthetic construct | AX589721.1 | GI:27901011 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |
| other sequences; artificial sequences. | AX589721.1 | GI:27901011 | Sequence 28 from Patent WO02081621. | 39 bp | DNA | linear | PAT 24-JAN-2003 |

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REFERENCE 1
AUTHORS      Loosmore,S.M. and Audonnet,J.C.
TITLE        Vaccine against the nile fever virus
JOURNAL      Patent: WO 02081621-A 28 17-OCT-2002;
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Qy 1 TCATGACTGCAATTCCGGTC 20
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Db 34 TCATGACTGCAATTCCGGTC 15

RESULT 3
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LOCUS          AX589711          33 bp      DNA      linear      PAT 24-JAN-2003
DEFINITION     Sequence 18 from Patent WO02081621.
ACCESSION      AX589711
VERSION        AX589711.1 GI:27901001
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE      1
AUTHORS        Loosmore,S.M. and Audonnet,J.C.
TITLE          Vaccine against the nile fever virus
JOURNAL        Patent: WO 02081621-A 18 17-OCT-2002;
              MERIAL (FR)
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Best Local Similarity 100.0%; Pred. No. 0.81;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCCGGT 19
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Db 28 TCATGACTGCAATTCCGGT 10

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LOCUS          AX589702          36 bp      DNA      linear      PAT 24-JAN-2003
DEFINITION     Sequence 9 from Patent WO02081621.
ACCESSION      AX589702
VERSION        AX589702.1 GI:27900992
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE      1
AUTHORS        Loosmore,S.M. and Audonnet,J.C.
TITLE          Vaccine against the nile fever virus
JOURNAL        Patent: WO 02081621-A 9 17-OCT-2002;
              MERIAL (FR)
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Qy 1 TCATGACTGCAATTCCGGT 19
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Db 28 TCATGACTGCAATTCCGGT 10

RESULT 5
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DEFINITION     Sequence 41 from Patent WO0160847.
ACCESSION      AX224249
VERSION        AX224249.1 GI:15554499
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE      1
AUTHORS        Kinney,R.M., Kinney,C.Y., Butrapet,S., Gubler,D.L. and
              Bhamarapravati,N.
TITLE          Avirulent, immunogenic flavivirus chimeras
JOURNAL        Patent: WO 0160847-A 41 23-AUG-2001;
              The Secretary, Department of Health and Human Services (US)
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Db 43 TCATGACTGCAATTCCGGT 25

RESULT 6
AX182204
LOCUS          AX182204          25 bp      DNA      linear      PAT 06-AUG-2001
DEFINITION     Sequence 14 from Patent WO0142441.
ACCESSION      AX182204
VERSION        AX182204.1 GI:15133479
KEYWORDS       .
SOURCE         synthetic construct
ORGANISM       other sequences; artificial sequences.
REFERENCE      1
AUTHORS        Reddy,S.I., Sadhu,L.I., Shukla,V.C. and Ferraiolo,G.I.
TITLE          Plasmid transformation
JOURNAL        Patent: WO 0142441-A 14 14-JUN-2001;
              International Centre for Genetic Engineering and Biotechnology (IT)
              Location/Qualifiers
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1 ATGACTGCAATT 12

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DEFINITION Sequence 17 from Patent WO0206497.
ACCESSION AX382013
VERSION AX382013.1 GI:19576835
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Reddy, V.S. and Sadhu, L.
TITLE Transplastonic plants
JOURNAL Patent: WO 0206497-A 17 24-JAN-2002;
JOURNAL International Centre for Genetic Engineering and Biotechnology (IT)
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RESULT 8
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DEFINITION JNK inhibitor.
ACCESSION BD173847
VERSION BD173847.1 GI:28415180
KEYWORDS WO 02062792-A/7.
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 34)
AUTHORS Okawa, S., Naruo, K., Miwatashi, S., Kimura, H. and Kawamoto, T.
TITLE JNK inhibitor
JOURNAL TAKEDA CHEMICAL INDUSTRIES LTD, SHIGENORI OKAWA, KENICHI NARUO, SEIJI MIWATASHI, HIROYUKI KIMURA, TOMOHIRO KAWAMOTO
COMMENT OS Artificial Sequence
PN WO 02062792-A/7
PD 15-AUG-2002
PF 01-FEB-2002 WO 2002JP000828
PR 02-FEB-2001 JP 01P 027570
PI SHIGENORI OKAWA, KENICHI NARUO, SEIJI MIWATASHI, HIROYUKI KIMURA,
PI TOMOHIRO KAWAMOTO
PC C07D417/04, C07D417/14, A61K31/4439, A61K31/4545, A61K31/506, PC
PC A61P43/00,
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ORIGIN
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Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 9 TCATGACTGCAA 20

RESULT 9
AR444320
LOCUS AR444320 51 bp DNA linear PAT 20-FEB-2004
DEFINITION Sequence 731 from patent US 6670464.
ACCESSION AR444320
VERSION AR444320.1 GI:42672099
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 51)
AUTHORS Shimkets, R.A. and Leach, M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: US 6670464-A 731 30-DEC-2003;
FEATURES Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GACTGCAATTCC 16
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Db 12 GACTGCAATTCC 23

RESULT 10
AR444321
LOCUS AR444321 51 bp DNA linear PAT 20-FEB-2004
DEFINITION Sequence 732 from patent US 6670464.
ACCESSION AR444321
VERSION AR444321.1 GI:42672100
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 51)
AUTHORS Shimkets, R.A. and Leach, M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: US 6670464-A 732 30-DEC-2003;
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source 1. .51
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Db 12 GACTGCAATTCC 23
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RESULT 11
LOCUS CQ536043 60 bp DNA linear PAT 30-JAN-2004
DEFINITION Sequence 5678 from Patent WO0210449.
ACCESSION CQ536043
VERSION CQ536043.1 GI:41502307
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Shoshan,A., Wasserman,A., Mintz E., Mintz L. and Faigler,S.
TITLE Oligonucleotide library for detecting rna transcripts and splice variants that populate a transcriptome
JOURNAL Patent: WO 0210449-A 5678 07-FEB-2002;
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Location/Qualifiers
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DEFINITION Poliovirus defective interfering particle 17 mRNA, partial cds.
ACCESSION M30219
VERSION M30219.1 GI:332915
KEYWORDS
SOURCE Poliovirus
ORGANISM Poliovirus
REFERENCE 1
AUTHORS Kuge,S., Saito,I. and Nomoto,A.
TITLE Primary structure of poliovirus defective-interfering particle genomes and possible generation mechanisms of the particles
JOURNAL J. Mol. Biol. 192 (3), 473-487 (1986)
MEDLINE 87169734
PUBMED 3031313
COMMENT Original source text: Poliovirus defective interfering particle 17, cDNA to viral RNA.
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Best Local Similarity 100.0%; Pred. No. 1.7e+04; Indels 0; Gaps 0;
Matches 12; Conservative 0; Mismatches 0;
Qy 5 GACTGCAATTCC 16
Db 33 GACTGCAATTCC 22
RESULT 13
LOCUS AR119503 15 bp DNA linear PAT 16-MAY-2001

DEFINITION Sequence 26 from patent US 6153382.
ACCESSION AR119503
VERSION AR119503.1 GI:14102202
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Karn,J., Gait,M.John., Heaphy,S. and Dingwall,C.
TITLE Viral growth inhibition
JOURNAL Patent: US 6153382-A 26 28-NOV-2000;
FEATURES
source 1..15
Location/Qualifiers
/organism="unknown"
/mol_type="unassigned DNA"
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Best Local Similarity 100.0%; Pred. No. 7.8e+04; Indels 0; Gaps 0;
Matches 11; Conservative 0; Mismatches 0;
Qy 11 AATTCGGTCT 21
Db 14 AATTCGGTCT 4
RESULT 14
LOCUS AR285636/c 16 bp RNA linear PAT 10-APR-2003
DEFINITION Sequence 8 from patent US 6528640.
ACCESSION AR285636
VERSION AR285636.1 GI:29723230
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 16)
AUTHORS Beigelman,L., Burgin,A., Beaudry,A., Karpeisky,A., Matulic-Adamic,J., Sweedler,D. and Zinnen,S.
TITLE Synthetic ribonucleic acids with RNase activity
JOURNAL Patent: US 6528640-A 8 04-MAR-2003;
FEATURES
source 1..16
Location/Qualifiers
/organism="unknown"
/mol_type="unassigned RNA"
ORIGIN
Query Match 47.8%; Score 11; DB 6; Length 16;
Best Local Similarity 100.0%; Pred. No. 7.8e+04; Indels 0; Gaps 0;
Matches 11; Conservative 0; Mismatches 0;
Qy 9 GCAATTCGGT 19
Db 13 GCAATTCGGT 3
RESULT 15
LOCUS AR397627/c 16 bp RNA linear PAT 18-DEC-2003
DEFINITION Sequence 8 from patent US 6617438.
ACCESSION AR397627
VERSION AR397627.1 GI:40134758
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 16)
AUTHORS Beigelman,L., Burgin,A.B., Beaudry,A., Karpeisky,A., Matulic-Adamic,J., Sweedler,D. and Zinnen,S.
TITLE Oligoribonucleotides with enzymatic activity
JOURNAL Patent: US 6617438-A 8 09-SEP-2003;
FEATURES
source 1..16
Location/Qualifiers

/organism="unknown"
/mol_type="unassigned RNA"

ORIGIN

Query Match 47.8%; Score 11; DB 6; Length 16;
Best Local Similarity 100.0%; Pred. No. 7.8e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 9 GCAATTCGGT 19
|||
Db 13 GCAATTCGGT 3

Search completed: September 6, 2005, 22:42:49
Job time : 1494 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: September 6, 2005, 20:30:00 ; Search time 233 Seconds
(without alignments)

584.352 Million cell updates/sec

Title: US-10-729-421-8

Perfect score: 23

Sequence: 1 tcatgactgaattccgtcttt 23

Scoring table: OLIGO_NUC

Gapop_60.0 , Gapext 60.0

Searched: 4390206 seqs, 2959870667 residues

Word size : 10

Total number of hits satisfying chosen parameters: 684

Minimum DB seq length: 0

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Post-processing: Listing first 45 summaries

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- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
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- 12: Geneseqn2004as:*
- 13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|-------|--------------------|
| 1 | 23 | 100.0 | 23 | 12 | ADQ30638 West Nile |
| 2 | 23 | 100.0 | 46 | 12 | ADQ30655 West Nile |
| C 3 | 20 | 87.0 | 36 | 8 | ABZ25451 PCR prime |
| C 4 | 20 | 87.0 | 36 | 9 | AAL55873 FC117 PCR |
| C 5 | 20 | 87.0 | 36 | 12 | ADM16871 Plasmid p |
| C 6 | 20 | 87.0 | 39 | 8 | ABZ25450 West Nile |
| C 7 | 20 | 87.0 | 39 | 9 | AAL55872 FC116 RT- |
| C 8 | 20 | 87.0 | 39 | 12 | ADM16870 West Nile |
| C 9 | 20 | 87.0 | 51 | 10 | ADCO6634 PCR prime |
| C 10 | 20 | 87.0 | 51 | 10 | ADCO6633 PCR prime |
| C 11 | 20 | 87.0 | 58 | 12 | ADM16878 Plasmid p |
| C 12 | 19 | 82.6 | 33 | 8 | ABZ25440 PCR prime |
| C 13 | 19 | 82.6 | 33 | 9 | AAL55862 FC110 PCR |
| C 14 | 19 | 82.6 | 33 | 12 | ADM16860 Plasmid p |
| C 15 | 19 | 82.6 | 36 | 8 | ABZ25431 West Nile |
| C 16 | 19 | 82.6 | 36 | 9 | AAL57662 CF107 RT- |
| C 17 | 19 | 82.6 | 36 | 12 | ADM16851 West Nile |
| C 18 | 19 | 82.6 | 51 | 10 | ADCO6635 PCR prime |
| C 19 | 19 | 82.6 | 56 | 4 | AAD14623 WN virus- |
| 20 | 17 | 73.9 | 17 | 6 | ACN09412 WN minus |

| | | | | | | |
|------|----|------|----|----|----------|--------------------|
| C 21 | 17 | 73.9 | 17 | 6 | ACN00067 | ACN00067 WN Hamme |
| C 22 | 17 | 73.9 | 17 | 6 | ACN13570 | ACN13570 WN minus |
| C 23 | 17 | 73.9 | 17 | 6 | ACN03509 | ACN03509 WN Zinzy |
| C 24 | 17 | 73.9 | 17 | 6 | ACN14150 | ACN14150 WN minus |
| C 25 | 17 | 73.9 | 17 | 6 | ACN05528 | ACN05528 WN Amber |
| C 26 | 17 | 73.9 | 17 | 6 | ACN12169 | ACN12169 WN minus |
| C 27 | 17 | 73.9 | 17 | 6 | ACN04748 | ACN04748 WN DNZY |
| C 28 | 17 | 73.9 | 17 | 6 | ACN09413 | ACN09413 WN minus |
| C 29 | 16 | 69.6 | 17 | 6 | ACN12168 | ACN12168 WN minus |
| C 30 | 16 | 69.6 | 17 | 6 | ACN05527 | ACN05527 WN Amber |
| C 31 | 16 | 69.6 | 17 | 6 | ACN12170 | ACN12170 WN minus |
| C 32 | 16 | 69.6 | 17 | 6 | ACN01482 | ACN01482 WN Inozv |
| C 33 | 16 | 69.6 | 30 | 10 | ADCO6605 | ADCO6605 WN 1st j |
| C 34 | 15 | 65.2 | 17 | 6 | ACN03510 | ACN03510 WN Zinzy |
| C 35 | 15 | 65.2 | 17 | 6 | ACN15245 | ACN15245 WN minus |
| C 36 | 15 | 65.2 | 17 | 6 | ACN01481 | ACN01481 WN Inozv |
| C 37 | 14 | 60.9 | 17 | 6 | ACN14149 | ACN14149 WN minus |
| C 38 | 14 | 60.9 | 17 | 6 | ACN13571 | ACN13571 WN Hamme |
| C 39 | 13 | 56.5 | 17 | 6 | ACN00068 | ACN00068 WN Hamme |
| C 40 | 13 | 56.5 | 17 | 6 | ACN04747 | ACN04747 WN DNZY |
| C 41 | 13 | 56.5 | 17 | 6 | ACN15244 | ACN15244 WN minus |
| C 42 | 13 | 56.5 | 25 | 9 | ACK26449 | ACK26449 Human mic |
| C 43 | 13 | 56.5 | 30 | 10 | ADCO6609 | ADCO6609 Chimeric |
| C 44 | 13 | 56.5 | 30 | 10 | ADCO6611 | ADCO6611 Chimeric |
| C 45 | 12 | 52.2 | 17 | 6 | ACN09414 | ACN09414 WN minus |

ALIGNMENTS

RESULT 1

ADQ30638
ID ADQ30638 standard; DNA; 23 BP.

AC ADQ30638;

DT 23-SEP-2004 (first entry)

DE West Nile Virus capture oligonucleotide WNVVC8.

ss; capture oligonucleotide; West Nile Virus; diagnosis.

OS West Nile virus.

PN WO2004055159-A2.

PD 01-JUL-2004.

PF 05-DEC-2003; 2003WO-US038750.

PR 12-DEC-2002; 2002US-0432850P.

PR 20-JUN-2003; 2003US-0480431P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2004-488058/46.

PT New isolated oligonucleotides for accurately diagnosing West Nile virus infection or for capturing, detecting and quantitating West Nile virus in blood samples.

PS Claim 1; SEQ ID NO 8; 56pp; English.

XX The invention relates to an isolated oligonucleotide not more than 60 nucleotides in length comprising a nucleotide sequence (S1) of at least 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g. CC 20, 21 or 23 bp) given in the specification derived from the West Nile Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence identity to the nucleotide sequence of (S1), or complements of (S1) and (S2). The oligonucleotide further comprises a detectable label at the 5'-end and/or the 3'-end. The detectable label is a fluorescent label

CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to a capture oligonucleotide of the invention.

SQ Sequence 23 BP; 4 A; 6 C; 4 G; 9 T; 0 U; 0 Other;

Query Match 100.0%; Score 23; DB 12; Length 23;
 Best Local Similarity 100.0%; Pred. No. 0.00091;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGTCTTT 23
 DB 1 TCATGACTGCAATTCGGTCTTT 23

RESULT 2

ADQ30655
 ID ADQ30655 standard; DNA; 46 BP.

AC ADQ30655;

XX 23-SEP-2004 (first entry)

XX West Nile Virus capture oligonucleotide poly-A-WNVVC8.

XX ss; capture oligonucleotide; West Nile Virus; diagnosis.

XX West Nile virus.

XX WO2004055159-A2.

XX 01-JUL-2004.

XX 05-DEC-2003; 2003WO-US038750.

XX 12-DEC-2002; 2002US-0432850P.

XX 20-JUN-2003; 2003US-0480431P.

XX (CHIR) CHIRON CORP.

XX Shyamala V;

XX WPI; 2004-488058/46.

XX New isolated oligonucleotides for accurately diagnosing West Nile virus
 PT infection or for capturing, detecting and quantitating West Nile virus in
 PT blood samples.

XX Example 1; SEQ ID NO 25; 56pp; English.

XX The invention relates to an isolated oligonucleotide not more than 60
 CC nucleotides in length comprising a nucleotide sequence (S1) of at least
 CC 10 contiguous nucleotides from any of the 28 nucleotide sequences (e.g.
 CC 20, 21 or 23 bp) given in the specification derived from the West Nile
 CC Virus (WNV) genome, a nucleotide sequence (S2) having 90% sequence
 CC identity to the nucleotide sequence of (S1), or complements of (S1) and
 CC (S2). The oligonucleotide further comprises a detectable label at the 5'-
 CC end and/or the 3'-end. The detectable label is a fluorescent label
 CC selected from 6-carboxyfluorescein (6-FAM), tetramethyl rhodamine
 CC (TAMRA), and 2',4',5',7'-tetrachloro-4-7-dichlorofluorescein (TET). The
 CC composition and methods are useful for accurately diagnosing West Nile
 CC virus infection or for capturing, detecting and quantitating West Nile
 CC virus in biological samples, particularly blood samples. This sequence
 CC corresponds to a capture oligonucleotide of the invention.

SQ Sequence 46 BP; 27 A; 6 C; 4 G; 9 T; 0 U; 0 Other;

Query Match 100.0%; Score 23; DB 12; Length 46;
 Best Local Similarity 100.0%; Pred. No. 0.00088;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGTCTTT 23
 DB 24 TCATGACTGCAATTCGGTCTTT 46

RESULT 3

ABZ25451/C
 ID ABZ25451 standard; DNA; 36 BP.

XX AC ABZ25451;

XX 27-MAR-2003 (first entry)

XX PCR primer FC117, SEQ ID 29.

XX Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus;
 KW WNV; PCR; primer; ss.

XX Synthetic.

XX WO200281621-A2.

XX 17-OCT-2002.

XX 05-APR-2002; 2002WO-FR001200.

XX 06-APR-2001; 2001FR-00004737.

XX (MERI-) MERIAL.

XX Loosmore SM, Audonnet JF;

XX WPI; 2003-111799/10.

XX Vaccine for treatment or prevention of West Nile virus (WNV) infection,
 PT for use in veterinary medicine, comprises a recombinant virus expressing
 PT a WNV structural protein.

XX Example 18; Page 41; 56pp; French.

XX The present invention relates to a vaccine for protecting horses, dogs,
 CC cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine
 CC comprises: (i) one or more recombinant avipox, NVVAC or MVA viruses that
 CC express one of the WNV proteins prM, M and E and (ii) a vehicle or
 CC excipient. The present sequence is a PCR primer, which was used in an
 CC example from the invention

XX Sequence 36 BP; 8 A; 7 C; 10 G; 11 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 8; Length 36;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCATGACTGCAATTCGGTCT 20
 DB 31 TCATGACTGCAATTCGGTCT 12

RESULT 4

AAL55873/C
 ID AAL55873 standard; DNA; 36 BP.

XX AC AAL55873;

XX 06-NOV-2003 (first entry)

XX FC117 PCR primer used to amplify the plasmid pFC115.

XX Immunogenic composition; West Nile fever virus; WNV; prM; M; membrane; E;
 KW pre-membrane protein; envelope; virucide; vaccine; FC117; primer; PCR;
 KW ss; plasmid pFC115.

```

OS Unidentified.
PN US2003104008-A1.
XX
XX
PD 05-JUN-2003.
XX
XX 04-APR-2002; 2002US-00116298.
XX
XX 06-APR-2001; 2001US-0281923P.
XX
XX (LOOS/) LOOSMORE S M.
PA (AUDO/) AUDONNET J F.
XX
XX Loosmore SM, Audonnet JF;
PI WPI; 2003-567944/53.
XX
XX New immunogenic composition comprising a recombinant avipox virus that
PT expresses in vivo in the animal the West Nile (WN) proteins prM, M or E,
PT useful for inducing an immunological response against WN virus.
XX
XX Example 18; Page 14; 24pp; English.
XX
XX The invention relates to a novel immunogenic composition for inducing an
CC immune response against West Nile fever virus (WNV) in an animal. The
CC composition comprises a vehicle or excipient and a recombinant avipox
CC virus that expresses in vivo in the animal the WNV proteins prM (pre-
CC membrane protein), M (membrane protein) or E (envelope protein). The
CC animal is selected from canine, feline, bovine, porcine, chicken, equine,
CC a duck, a goose or a turkey. The composition of the invention
CC demonstrates virucide activity and may be useful as a vaccine against
CC WNV. The current sequence is that of the FC117 PCR primer of the
CC invention which was used to amplify the plasmid pFC115
XX
XX Sequence 36 BP; 8 A; 7 C; 10 G; 11 T; 0 U; 0 Other;
SQ
Query Match 87.0%; Score 20; DB 9; Length 36;
Best Local Similarity 100.0%; Pred. No. 0.052;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCATGACTGCAATTCGGTC 20
DB 31 TCATGACTGCAATTCGGTC 12
RESULT 5
ADM16871/C
ID ADM16871 standard; DNA; 36 BP.
XX
XX ADM16871;
AC
XX
XX 20-MAY-2004 (first entry)
XX
XX Plasmid pFC115 PCR primer #1.
DE
XX
XX Immunogen; vaccine; West Nile virus; ss; PCR; primer.
XX
XX Synthetic.
OS
XX US2004037848-A1.
PN
XX
XX 26-FEB-2004.
XX
XX 26-FEB-2003; 2003US-00374953.
XX
XX 06-APR-2001; 2001US-0281923P.
PR
XX 04-APR-2002; 2002US-00116298.
XX
XX (AUDO/) AUDONNET J F.
PA (MINK/) MINK J M.
PA (LOOS/) LOOSMORE S M.
PA (KARA/) KARACA K.
XX

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PI Audonnet JF, Minke JM, Loosmore SM, Karaca K;
XX WPI; 2004-191012/18.
XX
XX Vaccine composition, useful in inducing an immune response against West
PT Nile virus, comprises a vector that contains heterologous nucleic acid
PT molecule(s), and that expresses in vivo in the animal a WNV protein.
XX
XX Example 18; SEQ ID NO 29; 36pp; English.
XX
XX The invention relates to an immunogenic or vaccine composition which
CC induces an immune response against West Nile virus (WNV) in an animal
CC susceptible to WNV comprises a vector that contains heterologous nucleic
CC acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV
CC prM and E; WNV M and E; WNV prM, WNV M and E, WNV polypeptide prM-E, WNV
CC polypeptide M-E, or WNV polypeptide prM-M-E. The composition is useful
CC for inducing an immunological or protective immune response against WNV
CC and against another pathogen of the animal. Also inducing an
CC immunological or protective immune response against WNV in an animal
CC comprises administering to the animal (a) the immunogenic or vaccine
CC composition and (b) a WNV isolated antigen, immunogen or epitope, where
CC (a) is administered prior to (b) in a prime-boost regimen, or (b) is
CC administered prior to (a) in a prime-boost regimen, or (a) and (b) are
CC administered together, either sequentially or in admixture. The present
CC sequence is used in the exemplification of the invention.
XX
XX Sequence 36 BP; 8 A; 7 C; 10 G; 11 T; 0 U; 0 Other;
SQ
Query Match 87.0%; Score 20; DB 12; Length 36;
Best Local Similarity 100.0%; Pred. No. 0.052;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCATGACTGCAATTCGGTC 20
DB 31 TCATGACTGCAATTCGGTC 12
RESULT 6
ABZ25450/C
ID ABZ25450 standard; DNA; 39 BP.
XX
XX ABZ25450;
AC
XX
XX 27-MAR-2003 (first entry)
DT
XX
XX West Nile Virus PCR primer FC116, SEQ ID 28.
DE
XX
XX Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus;
XX WNV; PCR; primer; ss.
XX
XX West Nile Virus.
OS
XX WO200281621-A2.
PN
XX
XX 17-OCT-2002.
PD
XX
XX 05-APR-2002; 2002WO-FR001200.
PF
XX
XX 06-APR-2001; 2001FR-00004737.
PR
XX
XX (MERI-) MERIAL.
PA
XX
XX Loosmore SM, Audonnet JF;
PI WPI; 2003-111799/10.
XX
XX Vaccine for treatment or prevention of West Nile virus (WNV) infection,
PT for use in veterinary medicine, comprises a recombinant virus expressing
PT a WNV structural protein.
XX
XX Example 17; Page 40; 56pp; French.
PS
XX
XX The present invention relates to a vaccine for protecting horses, dogs,
XX

```

CC cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine
 CC comprises: (i) one or more recombinant avipox, NYVAC or MVA viruses that
 CC express one of the WNV proteins prM, M and E and (ii) a vehicle or
 CC excipient. The present sequence is a PCR primer, which was used in an
 CC example from the invention

SQ Sequence 39 BP; 9 A; 6 C; 9 G; 15 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 8; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCCGGTC 20

Db 34 TCATGACTGCAATTCCGGTC 15

RESULT 7

AA155872/c
 ID AAL55872 standard; DNA; 39 BP.

XX AC AAL55872;

XX DT 06-NOV-2003 (first entry)

XX FC116 RT-PCR primer used to amplify West Nile fever virus RNA.

XX Immunogenic composition; West Nile fever virus; WNV; prM; M; membrane; E;
 KW pre-membrane protein; envelope; virucide; vaccine; FC116; RT-PCR; primer;
 KW PCR; ss.

XX OS Unidentified.

XX OS Synthetic.

XX FN US2003104008-A1.

XX PD 05-JUN-2003.

XX PF 04-APR-2002; 2002US-00116298.

XX PR 06-APR-2001; 2001US-0281923P.

XX PA (LOOS/) LOOSMORE S M.

XX PA (AUDO/) AUDONNET J F.

XX PI Loosmore SM, Audonnet JF;

XX WI WI; 2003-567944/53.

XX New immunogenic composition comprising a recombinant avipox virus that
 PT expresses in vivo in the animal the West Nile (WN) proteins prM, M or E,
 PT useful for inducing an immunological response against WN virus.

XX Example 17; Page 14; 24pp; English.

XX The invention relates to a novel immunogenic composition for inducing an
 CC immune response against West Nile fever virus (WNV) in an animal. The
 CC composition comprises a vehicle or excipient and a recombinant avipox
 CC virus that expresses in vivo in the animal the WNV proteins prM (pre-
 CC membrane protein), M (membrane protein) or E (envelope protein). The
 CC animal is selected from canine, feline, bovine, porcine, chicken, equine,
 CC a duck, a goose or a turkey. The composition of the invention
 CC demonstrates virucide activity and may be useful as a vaccine against
 CC WNV. The current sequence is that of the FC116 RT-PCR primer of the
 CC invention which was used to amplify West Nile fever virus RNA

SQ Sequence 39 BP; 9 A; 6 C; 9 G; 15 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 9; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCCGGTC 20

Db 34 TCATGACTGCAATTCCGGTC 15

RESULT 8

ADM16870/c
 ID ADM16870 standard; DNA; 39 BP.

XX AC ADM16870;

XX DT 20-MAY-2004 (first entry)

XX DE West Nile virus RT-PCR primer #10.

XX immunogen; vaccine; West Nile virus; ss; reverse transcriptase; RT-PCR;
 KW primer.

XX OS West Nile virus.

XX PN US2004037848-A1.

XX PD 26-FEB-2004.

XX PF 26-FEB-2003; 2003US-00374953.

XX PR 06-APR-2001; 2001US-0281923P.

XX PR 04-APR-2002; 2002US-00116298.

XX PA (AUDO/) AUDONNET J F.

XX PA (MINK/) MINK J M.

XX PA (LOOS/) LOOSMORE S M.

XX PA (KARA/) KARACA K.

XX PI Audonnet JF, Minke JM, Loosmore SM, Karaca K;

XX WI WI; 2004-191012/18.

XX Vaccine composition, useful in inducing an immune response against West
 PT Nile virus, comprises a vector that contains heterologous nucleic acid
 PT molecule(s), and that expresses in vivo in the animal a WNV protein.

XX Example 17; SEQ ID NO 28; 36pp; English.

XX The invention relates to an immunogenic or vaccine composition which
 CC induces an immune response against West Nile virus (WNV) in an animal
 CC susceptible to WNV comprises a vector that contains heterologous nucleic
 CC acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV
 CC prM and E; WNV M and E; WNV prM, WNV M and E, WNV polyprotein prM-E, WNV
 CC polyprotein M-E, or WNV polyprotein prM-M-E. The composition is useful
 CC for inducing an immunological or protective immune response against WNV
 CC and against another pathogen of the animal. Also inducing an
 CC immunological or protective immune response against WNV in an animal
 CC comprises administering to the animal (a) the immunogenic or vaccine
 CC composition and (b) a WNV isolated antigen, immunogen or epitope, where
 CC (a) is administered prior to (b) in a prime-boost regimen, or (b) is
 CC administered prior to (a) in a prime-boost regimen, or (a) and (b) are
 CC administered together, either sequentially or in admixture. The present
 CC sequence is used in the exemplification of the invention.

SQ Sequence 39 BP; 9 A; 6 C; 9 G; 15 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 12; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.052;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCCGGTC 20

Db 34 TCATGACTGCAATTCCGGTC 15

RESULT 9

ADC06634/c
 ID ADC06634 standard; DNA; 51 BP.

```
XX ADC06634;
XX AC
XX DT
XX DE 18-DEC-2003 (first entry)
XX DE PCR primer SEQ ID 35 used during construction of WNV/DEN4 chimeras.
XX KW West Nile virus; WNV; DEN4; Dengue virus type 4; virucide; vaccine; ss;
XX KW PCR; primer.
XX OS Unidentified.
XX PN WO2003059384-A1.
XX PD 24-JUL-2003.
XX PF 09-JAN-2003; 2003WO-US000594.
XX PR 10-JAN-2002; 2002US-0347281P.
XX PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX PI Pletnev AG, Putnak JR, Chanock RM, Murphy BR, Whitehead SS;
XX PI Blaney JE;
XX DR WPI; 2003-636686/60.
XX CC Novel nucleic acid chimera comprising nucleic acids encoding structural
XX PT protein from West Nile virus and non-structural proteins from wild-type
XX PT strain of dengue virus useful for producing live West Nile virus
XX PT vaccines.
XX PS Disclosure; Page 19; 53pp; English.
XX CC The invention relates to a novel nucleic acid chimera comprising a first
XX CC nucleotide sequence encoding at least one structural protein from a West
XX CC Nile virus (WNV) and a second nucleotide sequence encoding non-structural
XX CC proteins from a wild-type strain of Dengue virus (DEN), such as Dengue
XX CC virus type 4 (DEN4). The nucleotide of the invention demonstrates
XX CC virucide activity and may be useful for producing a WNV vaccine. The
XX CC current sequence is that of the PCR primer of the invention which was
XX CC used during the construction of the WNV/DEN4 chimeras.
XX SQ Sequence 51 BP; 20 A; 9 C; 13 G; 9 T; 0 U; 0 Other;
Query Match 87.0%; Score 20; DB 10; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.051;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCATGACTGCAATTCCGGTC 20
DB 46 TCATGACTGCAATTCCGGTC 27
RESULT 10
ADC06633/c
ID ADC06633 standard; DNA; 51 BP.
XX AC ADC06633;
XX DT
XX DE 18-DEC-2003 (first entry)
XX DE PCR primer SEQ ID 34 used during construction of WNV/DEN4 chimeras.
XX KW West Nile virus; WNV; DEN4; Dengue virus type 4; virucide; vaccine; ss;
XX KW PCR; primer.
XX OS Unidentified.
XX PN WO2003059384-A1.
XX PD 24-JUL-2003.
XX PF 09-JAN-2003; 2003WO-US000594.
XX PR 10-JAN-2002; 2002US-0347281P.
XX PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX PI Pletnev AG, Putnak JR, Chanock RM, Murphy BR, Whitehead SS;
XX PI Blaney JE;
XX DR WPI; 2003-636686/60.
XX CC Novel nucleic acid chimera comprising nucleic acids encoding structural
XX PT protein from West Nile virus and non-structural proteins from wild-type
XX PT strain of dengue virus useful for producing live West Nile virus
XX PT vaccines.
XX PS Disclosure; Page 20; 53pp; English.
XX CC The invention relates to a novel nucleic acid chimera comprising a first
XX CC nucleotide sequence encoding at least one structural protein from a West
XX CC Nile virus (WNV) and a second nucleotide sequence encoding non-structural
XX CC proteins from a wild-type strain of Dengue virus (DEN), such as Dengue
XX CC virus type 4 (DEN4). The nucleotide of the invention demonstrates
XX CC virucide activity and may be useful for producing a WNV vaccine. The
XX CC current sequence is that of the PCR primer of the invention which was
XX CC used during the construction of the WNV/DEN4 chimeras.
XX SQ Sequence 51 BP; 20 A; 9 C; 13 G; 9 T; 0 U; 0 Other;
Query Match 87.0%; Score 20; DB 10; Length 51;
Best Local Similarity 100.0%; Pred. No. 0.051;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCATGACTGCAATTCCGGTC 20
DB 46 TCATGACTGCAATTCCGGTC 27
RESULT 11
ADM16878/c
ID ADM16878 standard; DNA; 58 BP.
XX AC ADM16878;
XX DT
XX DE 20-MAY-2004 (first entry)
XX DE Plasmid pFC115 PCR primer #2.
XX KW immunogen; vaccine; West Nile virus; ss; PCR; primer.
XX OS Synthetic.
XX PN US2004037848-A1.
XX PD 26-FEB-2004.
XX PR 26-FEB-2003; 2003US-00374953.
XX PR 06-APR-2001; 2001US-0281923P.
XX PR 04-APR-2002; 2002US-00116298.
XX PA (AUDO/) AUDONNET J F.
XX PA (MINK/) MINK J M.
XX PA (LOOS/) LOOSMORE S M.
XX PA (KARA/) KARACA K.
XX PI Audonnet JF, Minke JM, Loosmore SM, Karaca K;
XX DR WPI; 2004-191012/18.
XX CC Vaccine composition, useful in inducing an immune response against West
XX PT Nile virus, comprises a vector that contains heterologous nucleic acid
XX PT molecule(s), and that expresses in vivo in the animal a WNV protein.
```

XX Example 28; SEQ ID NO 36; 36pp; English.

CC The invention relates to an immunogenic or vaccine composition which

CC induces an immune response against West Nile virus (WNV) in an animal

CC susceptible to WNV comprising a vector that contains heterologous nucleic

CC acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV

CC prM and E; WNV M and E; WNV prM, WNV M and E, WNV polyprotein prM-E, WNV

CC polyprotein M-E, or WNV polyprotein prM-M-E. The composition is useful

CC for inducing an immunological or protective immune response against WNV

CC and against another pathogen of the animal. Also inducing an

CC immunological or protective immune response against WNV in an animal

CC comprises administering to the animal (a) the immunogenic or vaccine

CC composition and (b) a WNV isolated antigen, immunogen or epitope, where

CC (a) is administered prior to (b) in a prime-boost regimen, or (b) is

CC administered prior to (a) in a prime-boost regimen, or (a) and (b) are

CC administered together, either sequentially or in admixture. The present

CC sequence is used in the exemplification of the invention.

XX SQ Sequence 58 BP; 13 A; 10 C; 14 G; 21 T; 0 U; 0 Other;

Query Match 87.0%; Score 20; DB 12; Length 58;

Best Local Similarity 100.0%; Pred. No. 0.051;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCCGGTC 20

Db 53 TCATGACTGCAATTCCGGTC 34

RESULT 12

AB225440/c

ID AB225440 standard; DNA; 33 BP.

XX AC AB225440;

XX DT 27-MAR-2003 (first entry)

XX DE PCR primer FC110, SEQ ID 18.

XX KW Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus;

XX KM WNV; PCR; primer; ss.

XX OS Synthetic.

XX PN WO200281621-A2.

XX PD 17-OCT-2002.

XX PF 05-APR-2002; 2002WO-FR001200.

XX PR 06-APR-2001; 2001FR-00004737.

XX PA (MERI-) MERIAL.

XX PI Loosmore SM, Audonnet JF;

XX DR WPI; 2003-111799/10.

XX Vaccine for treatment or prevention of West Nile virus (WNV) infection,

PT for use in veterinary medicine, comprises a recombinant virus expressing

PT a WNV structural protein.

XX Example 9; Page 34; 56pp; French.

XX The present invention relates to a vaccine for protecting horses, dogs,

CC cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine

CC comprises: (i) one or more recombinant avipox, NVVAC or MVA viruses that

CC express one of the WNV proteins prM, M and E and (ii) a vehicle or

CC excipient. The present sequence is a PCR primer, which was used in an

CC example from the invention

XX SQ Sequence 33 BP; 7 A; 7 C; 9 G; 10 T; 0 U; 0 Other;

Query Match 82.6%; Score 19; DB 8; Length 33;

Best Local Similarity 100.0%; Pred. No. 0.2;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCCGGT 19

Db 28 TCATGACTGCAATTCCGGT 10

RESULT 13

AAL55862/c

ID AAL55862 standard; DNA; 33 BP.

XX AC AAL55862;

XX DT 06-NOV-2003 (first entry)

XX DE FC110 PCR primer used to amplify the plasmid pFC105.

XX KW Immunogenic composition; West Nile fever virus; WNV; prM; M; membrane; E;

XX KM pre-membrane protein; envelope; virucide; vaccine; FC110; primer; PCR;

XX OS ss; plasmid pFC105.

XX PN Unidentified.

XX PD US2003104008-A1.

XX PF 05-JUN-2003.

XX PR 04-APR-2002; 2002US-00116298.

XX PR 06-APR-2001; 2001US-0281923P.

XX PA (LOOS/) LOOSMORE S M.

XX PA (AUDO/) AUDONNET J F.

XX PI Loosmore SM, Audonnet JF;

XX DR WPI; 2003-567944/53.

XX New immunogenic composition comprising a recombinant avipox virus that

PT expresses in vivo in the animal the West Nile (WN) proteins prM, M or E,

PT useful for inducing an immunological response against WN virus.

XX Example 10; Page 12; 24pp; English.

XX The invention relates to a novel immunogenic composition for inducing an

CC immune response against West Nile fever virus (WNV) in an animal. The

CC composition comprises a vehicle or excipient and a recombinant avipox

CC virus that expresses in vivo in the animal the WNV proteins prM (pre-

CC membrane protein), M (membrane protein) or E (envelope protein). The

CC animal is selected from canine, feline, bovine, porcine, chicken, equine,

CC a duck, a goose or a turkey. The composition of the invention

CC demonstrates virucide activity and may be useful as a vaccine against

CC WNV. The current sequence is that of the FC110 PCR primer of the

CC invention which was used to amplify the plasmid pFC105

XX SQ Sequence 33 BP; 7 A; 7 C; 9 G; 10 T; 0 U; 0 Other;

Query Match 82.6%; Score 19; DB 9; Length 33;

Best Local Similarity 100.0%; Pred. No. 0.2;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCCGGT 19

Db 28 TCATGACTGCAATTCCGGT 10

RESULT 14

ADM16860/c

ID ADM16860 standard; DNA; 33 BP.

XX


```

AC ADM16860;
XX
XX
DT 20-MAY-2004 (first entry)
XX
XX DE Plasmid pFC 105 PCR primer #1.
XX
XX KW immunogen; vaccine; West Nile virus; ss; PCR; primer.
XX
XX OS Synthetic.
XX
XX PN US2004037848-A1.
XX
XX PD 26-FEB-2004.
XX
XX PF 26-FEB-2003; 2003US-00374953.
XX
XX PR 06-APR-2001; 2001US-0281923P.
XX
XX PR 04-APR-2002; 2002US-00116298.
XX
XX PA (AUO//) AUDONNET J F.
XX
XX PA (MINK//) MINKE J M.
XX
XX PA (LOOS//) LOOSMORE S M.
XX
XX PA (KARA//) KARACA K.
XX
XX PI Audonnet JF, Minke JM, Loosmore SM, Karaca K;
XX
XX DR WPI; 2004-191012/18.
XX
XX PT Vaccine composition, useful in inducing an immune response against West
XX
XX PT Nile virus, comprises a vector that contains heterologous nucleic acid
XX
XX PT molecule(s), and that expresses in vivo in the animal a WNV protein.
XX
XX PS Example 10; SEQ ID NO 18; 36pp; English.
XX
XX CC The invention relates to an immunogenic or vaccine composition which
XX
XX CC induces an immune response against West Nile virus (WNV) in an animal
XX
XX CC susceptible to WNV comprises a vector that contains heterologous nucleic
XX
XX CC acid molecule(s) and that expresses in vivo in the animal a WNV E; WNV
XX
XX CC prM and E; WNV M and E; WNV prM, WNV M and E, WNV polyprotein prM-E, WNV
XX
XX CC polyprotein M-E, or WNV polyprotein prM-M-E. The composition is useful
XX
XX CC for inducing an immunological or protective immune response against WNV
XX
XX CC and against another pathogen of the animal. Also inducing an
XX
XX CC immunological or protective immune response against WNV in an animal
XX
XX CC comprises administering to the animal (a) the immunogenic or vaccine
XX
XX CC composition and (b) a WNV isolated antigen, immunogen or epitope, where
XX
XX CC (a) is administered prior to (b) in a prime-boost regimen, or (b) is
XX
XX CC administered prior to (a) in a prime-boost regimen, or (a) and (b) are
XX
XX CC administered together, either sequentially or in admixture. The present
XX
XX CC sequence is used in the exemplification of the invention.
XX
XX SQ Sequence 33 BP; 7 A; 7 C; 9 G; 10 T; 0 U; 0 Other;

Query Match 82.6%; Score 19; DB 12; Length 33;
Best Local Similarity 100.0%; Pred. No. 0.2;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCGGT 19
Db 28 TCATGACTGCAATTCGGT 10

RESULT 15
ABZ25431/C
ID ABZ25431 standard; DNA; 36 BP.
XX
XX AC ABZ25431;
XX
XX DT 27-MAR-2003 (first entry)
XX
XX DE West Nile Virus PCR primer FC107, SEQ ID 9.
XX
XX KW Virucide; vaccine; horse; dog; cat; cattle; pig; bird; West Nile virus;
XX
XX KW WNV; PCR; primer; ss.

```

```

XX
XX OS West Nile Virus.
XX
XX PN WO200281621-A2.
XX
XX PD 17-OCT-2002.
XX
XX PF 05-APR-2002; 2002WO-FR001200.
XX
XX PR 06-APR-2001; 2001FR-00004737.
XX
XX PA (MERI-) MERIAL.
XX
XX PI Loosmore SM, Audonnet JF;
XX
XX DR WPI; 2003-111799/10.
XX
XX PT Vaccine for treatment or prevention of West Nile virus (WNV) infection,
XX
XX PT for use in veterinary medicine, comprises a recombinant virus expressing
XX
XX PT a WNV structural protein.
XX
XX PS Example 7; Page 31; 56pp; French.
XX
XX CC The present invention relates to a vaccine for protecting horses, dogs,
XX
XX CC cats, cattle, pigs and birds against West Nile virus (WNV). The vaccine
XX
XX CC comprises: (i) one or more recombinant avipox, NYVAC or MVA viruses that
XX
XX CC express one of the WNV proteins prM, M and E and (ii) a vehicle or
XX
XX CC excipient. The present sequence is a PCR primer, which was used in an
XX
XX CC example from the invention
XX
XX SQ Sequence 36 BP; 8 A; 6 C; 8 G; 14 T; 0 U; 0 Other;

Query Match 82.6%; Score 19; DB 8; Length 36;
Best Local Similarity 100.0%; Pred. No. 0.2;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGACTGCAATTCGGT 19
Db 31 TCATGACTGCAATTCGGT 13

Search completed: September 6, 2005, 22:17:49
Job time : 234 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model
Run on: September 6, 2005, 21:56:10 ; Search time 1636 Seconds
(without alignments)
535.133 Million cell updates/sec

Title: US-10-729-421-8
Perfect score: 23
Sequence: 1 tcatgactgaattccggtcttt 23

Scoring table: OLIGO NUC
Gapop 60.0, Gapext 60.0

Searched: 34239544 seqs, 19032134700 residues
Word size : 10

Total number of hits satisfying chosen parameters: 61

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Listing first 45 summaries

Database : EST:
1: gb_est1:
2: gb_est2:
3: gb_hc:
4: gb_est3:
5: gb_est4:
6: gb_est5:
7: gb_est6:
8: gb_gse1:
9: gb_gse2:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|-----------|--------------------|
| C 1 | 12 | 52.2 | 55 | AA619888 | AA619888 vl58h07.8 |
| C 2 | 11 | 47.8 | 29 | CC456749 | CC456749 SALK 1002 |
| C 3 | 11 | 47.8 | 33 | AZ817376 | AZ817376 2M0086N22 |
| C 4 | 11 | 47.8 | 41 | BH791854 | BH791854 SALK 0618 |
| C 5 | 11 | 47.8 | 41 | BH812487 | BH812487 SALK 0618 |
| C 6 | 11 | 47.8 | 50 | BG153713 | BG153713 nag57906 |
| C 7 | 11 | 47.8 | 52 | AW249907 | AW249907 2821659.3 |
| C 8 | 11 | 47.8 | 55 | CN564600 | CN564600 tag20b04 |
| C 9 | 11 | 47.8 | 57 | BH918919 | BH918919 3526.1.63 |
| C 10 | 10 | 43.5 | 19 | AJ588850 | AJ588850 Arabidops |
| C 11 | 10 | 43.5 | 25 | AG197078 | AG197078 Pan trogl |
| C 12 | 10 | 43.5 | 34 | BX660142 | BX660142 Arabidops |
| C 13 | 10 | 43.5 | 36 | BH810737 | BH810737 SALK 0511 |
| C 14 | 10 | 43.5 | 36 | DMB546528 | AJ546528 Drosophi1 |
| C 15 | 10 | 43.5 | 37 | AA972482 | AA972482 op42d03.8 |
| C 16 | 10 | 43.5 | 37 | AA937582 | AA937582 wp81b11.x |
| C 17 | 10 | 43.5 | 37 | U44311 | U44311 ENU44311 As |
| C 18 | 10 | 43.5 | 37 | BX531847 | BX531847 Arabidops |
| C 19 | 10 | 43.5 | 39 | AV833097 | AV833097 AV833097 |
| C 20 | 10 | 43.5 | 39 | BX568339 | BX568339 BX568339 |
| C 21 | 10 | 43.5 | 39 | TA160803P | TA160803 T. brucei |
| C 22 | 10 | 43.5 | 41 | CR397281 | CR397281 Arabidops |
| C 23 | 10 | 43.5 | 42 | BX943895 | BX943895 Arabidops |
| C 24 | 10 | 43.5 | 44 | AL752437 | AL752437 Arabidops |

| | | | | | | |
|------|----|------|----|---|----------|--------------------|
| C 25 | 10 | 43.5 | 46 | 6 | CF049474 | CF049474 QCL37a04. |
| C 26 | 10 | 43.5 | 46 | 7 | CN753215 | CN753215 2M0086N22 |
| C 27 | 10 | 43.5 | 47 | 8 | BH000511 | BH000511 APHL3LD-X |
| C 28 | 10 | 43.5 | 48 | 9 | BG253356 | BG253356 602362952 |
| C 29 | 10 | 43.5 | 48 | 9 | BX945507 | BX945507 Arabidops |
| C 30 | 10 | 43.5 | 48 | 9 | CR356946 | CR356946 Arabidops |
| C 31 | 10 | 43.5 | 48 | 9 | CU528770 | CU528770 ASV9G01.f |
| C 32 | 10 | 43.5 | 49 | 5 | BQ100687 | BQ100687 1j22c04.x |
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| C 35 | 10 | 43.5 | 50 | 1 | AU103081 | AU103081 AU103081 |
| C 36 | 10 | 43.5 | 50 | 1 | AU103082 | AU103082 AU103082 |
| C 37 | 10 | 43.5 | 50 | 8 | BH612727 | BH612727 SALK_0331 |
| C 38 | 10 | 43.5 | 52 | 1 | AA068274 | AA068274 mm53C01.f |
| C 39 | 10 | 43.5 | 52 | 2 | BF632337 | BF632337 NF018E03D |
| C 40 | 10 | 43.5 | 52 | 8 | AZ629385 | AZ629385 1M0482N16 |
| C 41 | 10 | 43.5 | 52 | 9 | BX122966 | BX122966 Danio rer |
| C 42 | 10 | 43.5 | 53 | 4 | BG524434 | BG524434 42-53 Sfe |
| C 43 | 10 | 43.5 | 53 | 7 | CN870218 | CN870218 001204AAO |
| C 44 | 10 | 43.5 | 53 | 8 | BH252021 | BH252021 SALK 0124 |
| C 45 | 10 | 43.5 | 54 | 8 | AZ576149 | AZ576149 AST-T11C0 |

ALIGNMENTS

RESULT 1
AA619888/c 55 bp mRNA linear EST 09-OCT-1997
LOCUS vl58h07.s1 Knowles Solter mouse 2 cell Mus musculus CDNA clone
DEFINITION IMAGE:976477 5', mRNA sequence.

ACCESSION AA619888
VERSION AA619888.1 GI:2523764

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

REFERENCE 1 (bases 1 to 55)

AUTHORS

Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
Waterston, R.

TITLE The WashU-HHMI Mouse EST Project

JOURNAL Unpublished (1996)

COMMENT

Contact: Marra M/Mouse EST Project

WashU-HHMI Mouse EST Project

Washington University School of MedicineP

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: mouseest@wustl.edu

This clone is available royalty-free through LLNL ; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

MGI:557205.

FEATURES

source

1. 55
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J x DBA/2J F1"
/db_xref="taxon:10090"
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/lab_host="DH10B"
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/note="Organ: embryo; Vector: pBluescribe (modified);
Site 1: MluI; Site 2: SalI; Cloned unidirectionally from
mRNA prepared from 13,500 2-cell stage embryos. Primer:
SalI (dl): 5'-CGGTCGACCGTCGACCGTTTCTTTTCTTT-3', CDNAS
were cloned into the MluI/SalI sites of a modified
pBluescribe vector using commercial linkers (NEB).

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ORIGIN
Query Match          52.2%; Score 12; DB 1; Length 55;
Best Local Similarity 100.0%; Pred. No. 7.5e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GACTGCAATTCC 16
   |||||
DB 21 GACTGCAATTCC 10

RESULT 2
CC456749          29 bp DNA linear GSS 30-MAY-2003
LOCUS SALK_100255.25.15.x Arabidopsis thaliana TDNA insertion lines
DEFINITION Arabidopsis thaliana genomic clone SALK_100255.25.15.x, genomic
survey sequence.
ACCESSION CC456749
VERSION CC456749
KEYWORDS GSS
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana

REFERENCE
AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadrinac,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.
TITLE A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL Unpublished (2001)
COMMENT Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA. This sequence lies within 300 bases of the 5' end of
At2g27775.
Class: TDNA tagged.
Location/Qualifiers
1..29
/organism="Arabidopsis thaliana"
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/ecotype="Col-0"
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/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/clone="SALK_100255.25.15.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
Query Match          47.8%; Score 11; DB 8; Length 29;
Best Local Similarity 100.0%; Pred. No. 3.1e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 GCAATTCGGT 19
   |||||
DB 12 GCAATTCGGT 22

RESULT 3
AZ817376          33 bp DNA linear GSS 20-FEB-2001
LOCUS 2M0086N22R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
DEFINITION

clone UUGC2M0086N22 R, genomic survey sequence.
AZ817376          47.8%; Score 11; DB 8; Length 33;
Best Local Similarity 100.0%; Pred. No. 3.1e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TGACTGCAATT 14
   |||||
DB 23 TGACTGCAATT 33

RESULT 4
BH791854          41 bp DNA linear GSS 02-APR-2002
LOCUS BH791854
DEFINITION SALK_061833.40.05.x Arabidopsis thaliana TDNA insertion lines

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Arabidopsis thaliana genomic clone SALK_061833.40.05.x, genomic
survey sequence.
ACCESSION      BH791854
VERSION        BH791854.1  GI:19886147
KEYWORDS
SOURCE         Arabidopsis thaliana (thale cress)
ORGANISM       Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE      1 (bases 1 to 41)
AUTHORS        Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadrinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.
TITLE          A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL        Unpublished (2001)
COMMENT        Contact: Joseph R. Ecker
                Salk Institute Genomic Analysis Laboratory (SIGnAL)
                The Salk Institute for Biological Studies
                10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
                Tel: 858 453 4100 x1752
                Fax: 858 558 6379
                Email: eckers@salk.edu
                This is single pass sequence recovered from the left border of
                TDNA.
                Class: TDNA tagged.
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                1..41
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                /note="PCR was performed on Arabidopsis thaliana lines
                each of which contains one or more TDNA insertion
                elements. The resultant fragment for each line was
                directly sequenced to determine the genomic sequence at
                the site of insertion. Details of the protocols used can
                be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
Query Match      47.8%; Score 11; DB 8; Length 41;
Best Local Similarity 100.0%; Pred. No. 3.1e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      3 ATGACTGCAAT 13
        |||||
Db      38 ATGACTGCAAT 28

RESULT 5
BH812487/c
LOCUS      BH812487
DEFINITION SALK_061833 Arabidopsis thaliana TDNA insertion lines Arabidopsis
            thaliana genomic clone SALK_061833, genomic survey sequence.
ACCESSION  BH812487
VERSION     BH812487.1  GI:20390942
KEYWORDS
SOURCE      Arabidopsis thaliana (thale cress)
ORGANISM    Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE    1 (bases 1 to 41)
AUTHORS      Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadrinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.
TITLE        A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL      Unpublished (2001)
COMMENT      Contact: Joseph R. Ecker

Arabidopsis thaliana genomic clone SALK_061833.40.05.x, genomic
survey sequence.
ACCESSION      BH791854
VERSION        BH791854.1  GI:19886147
KEYWORDS
SOURCE         Arabidopsis thaliana (thale cress)
ORGANISM       Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE      1 (bases 1 to 41)
AUTHORS        Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R.,
Gadrinab,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L.,
Shinn,P., Zimmerman,J. and Ecker,J.R.
TITLE          A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL        Unpublished (2001)
COMMENT        Contact: Joseph R. Ecker
                Salk Institute Genomic Analysis Laboratory (SIGnAL)
                The Salk Institute for Biological Studies
                10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
                Tel: 858 453 4100 x1752
                Fax: 858 558 6379
                Email: eckers@salk.edu
                This is single pass sequence recovered from the left border of
                TDNA.
                Class: TDNA tagged.
                Location/Qualifiers
                1..41
                /organism="Arabidopsis thaliana"
                /mol_type="genomic DNA"
                /ecotype="Col-0"
                /db_xref="taxon:3702"
                /clone="SALK_061833.40.05.x"
                /note="PCR was performed on Arabidopsis thaliana lines
                each of which contains one or more TDNA insertion
                elements. The resultant fragment for each line was
                directly sequenced to determine the genomic sequence at
                the site of insertion. Details of the protocols used can
                be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
Query Match      47.8%; Score 11; DB 8; Length 41;
Best Local Similarity 100.0%; Pred. No. 3.1e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      3 ATGACTGCAAT 13
        |||||
Db      38 ATGACTGCAAT 28

RESULT 6
BG153713/c
LOCUS      BG153713
DEFINITION nags57g06.x1 NCI_CGAP_Co26 Homo sapiens cDNA clone IMAGE:4225738 3',
            mRNA sequence.
ACCESSION  BG153713
VERSION     BG153713.1  GI:12665743
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE    1 (bases 1 to 50)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
            Unpublished (1997)
            Contact: Robert Strausberg, Ph.D.
            Email: cgaps-r@mail.nih.gov
            CDNA Library Preparation: David B. Krizman, Ph.D.
            CDNA Library Arrayed by: The I.M.A.G.E. Consortium/LLNL
            Cloning by: Washington University Genome Sequencing Center
            Cloning Distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL, send email to:
            infoimage.llnl.gov
            Seq primer: -40UP from Gibco.
            Location/Qualifiers
            1..50
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="IMAGE:4225738"
            /tissue_type="normal colonic mucosa"
            /lab_host="DH10B"
            /clone_lib="NCI CGAP Co26"
            /notes="Organ: colon; Vector: pAMP1; mRNA made from normal
            colonic mucosa, cDNA made by oligo-dT priming.
            Directionally cloned into UDG sites. Size-selected on
            agarose gel, average insert size 300 bp. Primary library."

```

```

Salk Institute Genomic Analysis Laboratory (SIGnAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: eckers@salk.edu
This is single pass sequence recovered from the left border of
TDNA.
Class: TDNA tagged.
Location/Qualifiers
1..41
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Col-0"
/db_xref="taxon:3702"
/clone="SALK_061833"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

ORIGIN
Query Match      47.8%; Score 11; DB 8; Length 41;
Best Local Similarity 100.0%; Pred. No. 3.1e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      3 ATGACTGCAAT 13
        |||||
Db      38 ATGACTGCAAT 28

RESULT 6
BG153713/c
LOCUS      BG153713
DEFINITION nags57g06.x1 NCI_CGAP_Co26 Homo sapiens cDNA clone IMAGE:4225738 3',
            mRNA sequence.
ACCESSION  BG153713
VERSION     BG153713.1  GI:12665743
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE    1 (bases 1 to 50)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
            Unpublished (1997)
            Contact: Robert Strausberg, Ph.D.
            Email: cgaps-r@mail.nih.gov
            CDNA Library Preparation: David B. Krizman, Ph.D.
            CDNA Library Arrayed by: The I.M.A.G.E. Consortium/LLNL
            Cloning by: Washington University Genome Sequencing Center
            Cloning Distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL, send email to:
            infoimage.llnl.gov
            Seq primer: -40UP from Gibco.
            Location/Qualifiers
            1..50
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="IMAGE:4225738"
            /tissue_type="normal colonic mucosa"
            /lab_host="DH10B"
            /clone_lib="NCI CGAP Co26"
            /notes="Organ: colon; Vector: pAMP1; mRNA made from normal
            colonic mucosa, cDNA made by oligo-dT priming.
            Directionally cloned into UDG sites. Size-selected on
            agarose gel, average insert size 300 bp. Primary library."

```

CDNA Library Preparation: David B. Krizman, Ph.D.
Reference: Krizman et al. (1996) Cancer Research
56:5380-5393."

ORIGIN

Query Match 47.8%; Score 11; DB 4; Length 50;
Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CATGACTGCAA 12
|||||
DB 13 CATGACTGCAA 3

RESULT 7

AW249907/c
LOCUS AW249907 52 bp mRNA linear EST 07-JAN-2000
DEFINITION 2821659.3prime NIH_MGC_7 Homo sapiens cDNA clone IMAGE:2821659 3',
mRNA sequence.
ACCESSION AW249907
VERSION AW249907.1 GI:6592900
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 52)
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
TITLE NIH-MGC http://mgc.nci.nih.gov/.
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)
COMMENT Unpublished (1999)
Other ESTs: 2821659.5prime
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov

Tissue Procurement: DCTP/DRP CDNA Library Preparation: Ling
Hong/Rubin Laboratory cDNA Library Arrayed by: The I.M.A.G.E.
Consortium (LNL) DNA Sequencing by: Berkeley MGC sequencing
project Clone distribution: MGC clone distribution information can
be found through the I.M.A.G.E. Consortium/LNL at:
www.bio.lnl.gov/bbrp/image.html Base Calling / Quality
Scores: PHRED from University of Washington Genome Center. Vector
Trimming: crosses match from University of Washington Genome Center
PHRAP suite. Poly-T Identification: patMatch.pl from Berkeley
Drosophila Genome Project. University of Washington Genome Center:
http://www.genome.washington.edu Low Quality Sequence: 30
contiguous PHRED high quality bases following vector sequence. Very
Low Quality Sequence: Trace file contained 52 contiguous distinct
peaks following vector sequence. Polyadenylation: Based upon the
presence of a XhoI site followed by a run of 14 or more T residues
at the beginning of the sequence, this cDNA insert was
polyadenylated.

Plate: L16W7 row: G column: 4

High quality sequence stop: 30.

FEATURES

source
location/Qualifiers
1..52
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2821659"
/tissue_type="small cell carcinoma"
/cell_line="MGC3"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC 7"
/note="Organ: lung; Vector: pOT87; Site 1: XhoI; Site 2:
EcoRI; cDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGACGAG(G). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."

ORIGIN

Query Match 47.8%; Score 11; DB 2; Length 52;

Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ACTGCAATTC 16
|||||
DB 41 ACTGCAATTC 31

RESULT 8

CNS564600/c
LOCUS CNS564600 55 bp mRNA linear EST 03-MAY-2004
DEFINITION tag20B04.x1 Hydra EST -Kiel 1 Hydra magnipapillata cDNA 3', mRNA
sequence.
ACCESSION CNS564600.1 GI:46973904
VERSION CNS564600
KEYWORDS EST.
SOURCE Hydra magnipapillata
ORGANISM Eukaryota; Metazoa; Cnidaria; Hydrozoa; Hydroida; Anthomedusae;
Hydridae; Hydra.
REFERENCE 1 (bases 1 to 55)
AUTHORS Bode,H., Blumberg,B., Steele,R., Wigge,P., Gee,L., Nguyen,Q.,
Martinez,D., Kibler,D., Hampson,S., Clifton,S., Pape,D., Marra,M.,
Hillier,L., Martin,J., Wylie,T., Dente,M., Theising,B., Bowers,Y.,
Gibbons,M., Ritter,E., Bennett,J., Ronko,I., Tsagarishvili,R.,
Maguire,L., Kennedy,S., Waterston,R. and Wilson,R.
TITLE WashU Hydra EST Project
JOURNAL Unpublished (2002)
COMMENT Contact: Hans Bode
WashU Hydra EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu

Library was constructed by Konstantin Khalturin, Zoologisches
Institut, Univ. Kiel, Germany Library materials provided by Thomas
Bosch, Zoologisches Institut, CAU Kiel, Germany DNA sequencing by:
Washington University Genome Sequencing Center For information on
obtaining a clone please contact: Hans Bode (hrobe@uci.edu)
seq primer: degenerate primer.
Location/Qualifiers
1..55
/organism="Hydra magnipapillata"
/mol_type="mRNA"
/strain="105"
/db_xref="taxon:6085"
/lab_host="DH5a"
/clone_lib="Hydra EST -Kiel 1"
/note="Vector: pSPORT1; Site 1: Not I; Site 2: Sal I;
pSPORT 1 Vector is ampicillin resistant, M13 reverse
primer was used by us for sequencing of 5' parts of
inserts; 3' parts of cDNAs contain long polyA tracks which
makes sequencing from 3' direction complicated"

FEATURES

source
location/Qualifiers
47.8%; Score 11; DB 7; Length 55;
Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 TTCGGGCTTT 23
|||||
DB 37 TTCGGGCTTT 27

ORIGIN

Query Match 47.8%; Score 11; DB 7; Length 55;
Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 TTCGGGCTTT 23
|||||
DB 37 TTCGGGCTTT 27

RESULT 9
BH918919/c
LOCUS BH918919 57 bp DNA linear GSS 12-SEP-2002
DEFINITION 3526.1.63.1.A10.2EL.x.1 3526 - RescueMu Grid K Zea mays genomic,
genomic survey sequence.
ACCESSION BH918919
VERSION BH918919.1 GI:22808353
KEYWORDS GSS.

SOURCE ORGANISM
Zea mays
Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.

REFERENCE
1 (bases 1 to 57)

AUTHORS
Walbot,V.

TITLE
Maize genomic sequences found using engineered RescueMu transposon

JOURNAL
Unpublished (2001)

COMMENT
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Possible ligation site of ends cut by 2 different endonucleases.
Reverse complemented post-ligation sequence from source sequence.
Plate: 3526_1_63_1 row: 9
Class: transposon-tagged.

FEATURES
source
1..57
/organism="Zea mays"
/mol_type="genomic DNA"
/cultivar="mixed background W23/A188/B73"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="3526 - RescueMu Grid K"
/note="Organ: leaf; Vector: RescueMu (engineered from pBlueScript backbone); Site 1: BamHI; Site 2: BglII; RescueMu is a 4.9 kb, modified maize Mu transposon designed to allow plasmid rescue from total genomic DNA. Mu elements insert preferentially into transcription units. For more information on RescueMu, go to the web site 'www.zmdb.iastate.edu' and follow the links for 'RescueMu.' Grid K was grown at Molokai, Hawaii in winter 2000-2001. DNA was extracted from leaf punches, double digested using BamHI and BglII, and ligated to form circular plasmids. DH10B cells were transformed and then screened on LB plates with ampicillin."

ORIGIN
Query Match 47.8%; Score 11; DB 8; Length 57;
Best Local Similarity 100.0%; Pred. No. 3.2e+04;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CATGACTGCAA 12
|||||
Db 51 CATGACTGCAA 41

RESULT 10
AJ588850
LOCUS Arabidopsis thaliana T-DNA flanking sequence, left border, clone 19 bp DNA linear GSS 15-JAN-2004

DEFINITION 358F07 genomic survey sequence.

ACCESSION AJ588850

VERSION AJ588850.1 GI:37938474

KEYWORDS GSS; left border; T-DNA flanking sequence.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE
1
AUTHORS Brunaud,V., Balzergue,S., Dubreucq,B., Aubourg,S., Samson,F., Chauvin,S., Bechtold,N., Cruaud,C., DeRose,R., Pelletier,G., Lepoint,L., Caboche,M. and Lecharny,A.
TITLE T-DNA integration into the Arabidopsis genome depends on sequences of pre-insertion sites

JOURNAL EMBO Rep. 3 (12), 1152-1157 (2002)

MEDLINE 22363535
PUBMED 12446565
REFERENCE 2 (bases 1 to 19)
AUTHORS Balzergue,S.
TITLE Direct Submission
JOURNAL Submitted (23-OCT-2003) Balzergue S., UMRGV, INRA/CNRS, 2 rue PCR was performed on DNA from transformants of Arabidopsis thaliana plants from INRA (Versailles). The DNA fragment(s) resulting from the PCR were directly sequenced from the left or the right border to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed. Information to order the corresponding mutant line and a link to a database providing a graphical display of the insertion site are available at <http://dbgap.versailles.inra.fr/publiclines/>. This sequence has been generated in the framework of the French plant genomics program 'Genoplante' (<http://www.genoplante.com> and <http://genoplante-info.inbio.gen.fr>).

FEATURES
source
1..19
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultivar="Wassillewskija"
/db_xref="taxon:3702"
/clone_lib="Arabidopsis thaliana T-DNA insertion lines"
misc_feature 1..19
/note="T-DNA flanking sequence left border"

ORIGIN
Query Match 43.5%; Score 10; DB 9; Length 19;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 TGACTGCAAT 13
|||||
Db 3 TGACTGCAAT 12

RESULT 11
AG197078
LOCUS Pan troglodytes DNA, clone: RP43-077A10.TJ, genomic survey sequence. 25 bp DNA linear GSS 06-MAR-2004

DEFINITION AG197078

ACCESSION AG197078

VERSION AG197078.1 GI:45229254

KEYWORDS GSS.

SOURCE Pan troglodytes (chimpanzee)

ORGANISM Pan troglodytes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.

REFERENCE
1
AUTHORS Park,H., Kim,Y., Kim,S., Han,Y., Woo,T., Park,K., Eun,C.J., Hoon,S.T., Chu,M., Kim,H., Joo,S., Kim,C., Song,W. and Yoo,H.

TITLE BAC end sequences of Library RP-43

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 25)

AUTHORS Park,H., Kim,Y., Kim,S., Han,Y., Woo,T., Park,K., Eun,C.J., Hoon,S.T., Chu,M., Kim,H., Joo,S., Kim,C., Song,W. and Yoo,H.

TITLE Direct Submission

JOURNAL Submitted (07-JAN-2002) Hong-Seog Park, Korea Research Institute of Bioscience and Biotechnology (KRIBB), Genome Research Center (GRC); 52, Oun-dong, Yusong-gu, Daejeon 305-333, Korea (E-mail:redstone@mail.krribb.re.kr, URL:<http://phs.grc.krribb.re.kr/>, Tel:82-42-866-7181, Fax:82-42-860-4409)

COMMENT Clones are derived from the chimpanzee BAC library RP-43 This BAC end was generated during the R&D process and may have higher chance of clone tracking errors.
PRIMERS
Sequencing: TJ
LIBRARY
Vector : pBACe3.6

R.Site 1 : EcoRI
R.Site 2 : EcoRI.

FEATURES
source Location/Qualifiers

1..25
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="RP43-077A10.TJ"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RP-43 Chimpanzee Male BAC Library"

ORIGIN

Query Match 43.5%; Score 10; DB 9; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 CTGCAATTCC 16
|||||
Db 1 CTGCAATTCC 10

RESULT 12
LOCUS BX660142 34 bp DNA linear GSS 04-APR-2004
DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-650H01-021296,
genomic survey sequence.

ACCESSION BX660142
VERSION BX660142.1 GI:37616530

KEYWORDS

SOURCE GSS.

ORGANISM Arabidopsis thaliana (thale cress)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi

REFERENCE

1 Li, Y., Rosso, M.G., Strizhov, N., Viehoveer, P., and Weisshaar, B.

GABI-Kat SimpleSearch: a flanking sequence tag (FST) database for
the identification of T-DNA insertion mutants in Arabidopsis

thaliana

Bioinformatics 19 (11), 1441-1442 (2003)

2755829

12874060

REFERENCE

2 Rosso, M.G., Li, Y., Strizhov, N., Reiss, B., Dekker, K. and

Weisshaar, B.

An Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
flanking sequence tag-based reverse genetics

Plant Mol. Biol. 53 (1-2), 247-259 (2003)

23117147

14756321

REFERENCE

3 Strizhov, N., Li, Y., Rosso, M.G., Viehoveer, P., Dekker, K.A. and

Weisshaar, B.

High-throughput generation of sequence indexes from T-DNA

mutagenized Arabidopsis thaliana lines

BioTechniques 35 (6), 1164-1168 (2003)

14682050

REFERENCE

4 (bases 1 to 34)

Li, Y., Strizhov, N., Rosso, M.G. and Weisshaar, B.

Direct Submission

Submitted (31-MAR-2004) Weisshaar B., Max-Planck-Institut fuer

Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

This sequence has been recovered from the left border of the T-DNA.

It indicates an insertion within the locus defined by BAC Clone

T1923. Details on the protocols used for generation of the

sequence are described in References 1-3. The sequences are

generated at the MPI for Plant Breeding Research in the context of

the GABI-Kat project. GABI-Kat is part of the German Plant Genomics

program designated 'GABI'. Information on line availability can be

found at: <http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

FEATURES
source Location/Qualifiers

1..34

/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-650H01-021296"
/clone_lib="Arabidopsis thaliana T-DNA insertion lines"
/ecotype="Col-0"
/note="PCR was performed on DNA from Arabidopsis thaliana
plants (Ti) which were transformed with the T-DNA from
vector pAC161 (GenBank accession number: AJ537514). The
lines contain one or more T-DNA insertions. The DNA
fragment(s) resulting from the PCR were directly sequenced
to determine the genomic sequence flanking the insertion.
T-DNA derived sequences were removed."

ORIGIN

Query Match 43.5%; Score 10; DB 9; Length 34;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TGACTGCAAT 13
|||||
Db 17 TGACTGCAAT 26

RESULT 13

BH810737/c

LOCUS BH810737

DEFINITION

thaliana genomic clone SALK_051126, genomic survey sequence.

ACCESSION BH810737

VERSION BH810737.1 GI:20388555

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi

1 (bases 1 to 36)

Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R.,

Gadrinab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L.,

Shinn, P., Zimmerman, J. and Ecker, J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished (2001)

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGNAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@alk.edu

This is single pass sequence recovered from the left border of

TDNA.

Class: TDNA tagged.

Location/Qualifiers

1..36

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/ecotype="Col-0"

/db_xref="taxon:3702"

/clone="SALK_051126"

/note="PCR was performed on Arabidopsis thaliana TDNA insertion lines"

each of which contains one or more TDNA insertion

elements. The resultant fragment for each line was

directly sequenced to determine the genomic sequence at

the site of insertion. Details of the protocols used can

be found at http://signal.salk.edu/tdna_protocols.html

TDNA.

Class: TDNA tagged.

Location/Qualifiers

1..36

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/ecotype="Col-0"

/db_xref="taxon:3702"

/clone="SALK_051126"

/note="PCR was performed on Arabidopsis thaliana TDNA insertion lines"

each of which contains one or more TDNA insertion

elements. The resultant fragment for each line was

directly sequenced to determine the genomic sequence at

the site of insertion. Details of the protocols used can

be found at http://signal.salk.edu/tdna_protocols.html

TDNA.

Class: TDNA tagged.

Location/Qualifiers


```

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 14 TCCGGTCTTT 23
    |||||
Db 18 TCCGGTCTTT 9

RESULT 14
DME546528/c
LOCUS
DEFINITION
  Drosophila melanogaster flanking sequence of RS P element insertion
  P{RS5}5-HA-1904, Clone library P{RS5}, genomic survey sequence.
ACCESSION
  AJ546528
VERSION
  AJ546528.1 GI:28554603
KEYWORDS
  GSS; genome survey sequence.
SOURCE
  Drosophila melanogaster (fruit fly)
ORGANISM
  Drosophila melanogaster
  Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;
  Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
  Ephydroidea; Drosophilidae; Drosophila.

REFERENCE
1
AUTHORS
  Ryder,E.J., Ashburner,M., Bagunya,J., Blows,F., Bucheton,A.,
  Coulson,D., Dickson,B., Drummond,J., Glover,D., Gunton,N.,
  Hafen,B., Hall,S., Heisenberg,M., Lepesant,J.A., Maroy,P.,
  Mechler,B., O'Kane,C., Pflugfelder,G., Rasmuson-Lestander,A.,
  Reuter,G., Roote,J., Szidonya,J., Wang,S., Webster,J. and
  Russell,S.
TITLE
  Mapping of RS P element insertions in Drosophila melanogaster for
  the DrosDel second generation deficiency kit
JOURNAL
  Unpublished
REFERENCE
2 (bases 1 to 36)
AUTHORS
  Ryder,E.J.
TITLE
  Direct Submission
JOURNAL
  Submitted (17-FEB-2003) Ryder E.J., Department of Genetics,
  University of Cambridge, Downing Street, CB2 3EH, UNITED KINGDOM
COMMENT
  The insertion point of the P element is before base 1 of the
  sequence. Further information about this P element insertion line
  can be found at http://www.flyseq.org.uk and
  http://www.drosdel.org.uk.

FEATURES
. source
  1..36
  /organism="Drosophila melanogaster"
  /mol_type="genomic DNA"
  /db_xref="taxon:7227"
  /chromosome="2L"
  /clone="P{RS5}5-HA-1904"
  /clone_lib="P{RS5}"
  /note="read=5' end"
  misc_feature
  1..36
  /note="P element insertion in the 3' to 5' orientation"

ORIGIN
Query Match 43.5%; Score 10; DB 9; Length 36;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 ACTGCAATTC 15
    |||||
Db 23 ACTGCAATTC 14

RESULT 15
AA972482
LOCUS
DEFINITION
  op2d03.s1 Soares NPL T_GBC_S1 Homo sapiens cDNA clone
  IMAGE:1579493 3' similar to TR:Q13526 PIN1. ;, mRNA
  sequence.
ACCESSION
  AA972482
VERSION
  AA972482.1 GI:3145246
KEYWORDS
  EST.
SOURCE
  Homo sapiens (human)
ORGANISM
  Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

```

```

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 37)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 523 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
  1..37
  source
  /organism="Homo sapiens"
  /mol_type="mRNA"
  /db_xref="taxon:9606"
  /clone="IMAGE:1579493"
  /lab_host="DH10B"
  /clone_lib="Soares NPL T_GBC_S1"
  /note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
  a modified polylinker; Site_1: Not 1; Site_2: Eco RI;
  Equal amounts of plasmid DNA from three normalized
  libraries (fetal lung NBHL19W, testis NHT, and B-cell
  NCI CGAP GCBI) were mixed, and ss circles were made in
  vitro. Following HAP purification, this DNA was used as
  tracer in a subtractive hybridization reaction. The driver
  was PCR-amplified cDNAs from pools of 5,000 clones made
  from the same 3 libraries. The pools consisted of
  I.M.A.G.E. clones 297480-302087, 682632-687239,
  726408-728711, and 729096-731399. Subtraction by Bento
  Soares and M. Fatima Bonaldo."

ORIGIN
Query Match 43.5%; Score 10; DB 1; Length 37;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CATGACTGCA 11
    |||||
Db 1 CATGACTGCA 10

Search completed: September 6, 2005, 23:10:13
Job time : 1639 secs

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